

Substituted Quinoline CCR5 Receptor Antagonists

This application claims priority to U.S. Provisional Application Serial No. 60/451,687 filed June 27, 2002 the entirety of which is incorporated herein by reference.

BACKGROUND OF THE INVENTION

Chemoattractant cytokines or chemokines are a family of proinflammatory mediators that promote recruitment and activation of leukocytes (e.g., monocytes, 10 lymphocytes, and granulocytes) and also have additional effects on diverse cell types. They can be released by many kinds of tissue cells after activation. Continuous release of chemokines at sites of inflammation mediates the ongoing migration of effector cells in chronic inflammation. The chemokines characterized to date are related in primary structure. They share four conserved cysteines, which form disulfide 15 bonds. Based upon this conserved cysteine motif, the family is divided into two main branches, designated as the C-X-C chemokines (α -chemokines), and the C-C chemokines (β -chemokines), in which the first two conserved cysteines are separated by an intervening residue, or adjacent respectively (Baggiolini, M. and Dahinden, C. A., *Immunology Today*, 15:127-133 (1994)).

20 RANTES (Regulated on Activation, Normal T Expressed and Secreted), the macrophage inflammatory proteins 1 α . and 1 β (MIP-1 α . and MIP-1 β), and human monocyte chemotactic proteins 1-3 (MCP-1, MCP-2, MCP-3) are members of the C-C chemokine family and have been characterized as chemoattractants and activators of 25 monocytes or lymphocytes. Migration of leukocytes from blood vessels into diseased tissues is important to the initiation of normal disease-fighting inflammatory responses. This process, known as leukocyte recruitment, is also involved in the onset and progression of debilitating and life-threatening chronic inflammatory, allergic 30 inflammatory and autoimmune diseases. Compounds which block leukocyte recruitment to target tissues in inflammatory and autoimmune disease would be a highly effective therapeutic intervention. Abnormal production of chemokines, such as RANTES and MIP-1 α have been implicated in a wide range of human acute and chronic inflammatory diseases, including rheumatoid arthritis, multiple sclerosis, inflammatory bowel disease, endometriosis and respiratory diseases, such as asthma

and allergic disorders. A review of the role of chemokines in allergic inflammation is provided by Kita, H., et al., J. Exp. Med. 183, 2421-2426 (1996). Chemokines and their receptors have been implicated in the pathogenesis of a number of neurological disorders including ischemic stroke, trauma, Alzheimer's disease, multiple sclerosis
5 and cancer.

The chemokine receptors are members of a superfamily of G protein-coupled receptors (GPCR) which share structural features that reflect a common mechanism of action of signal transduction (Gerard, C. and Gerard, N. P., Annu Rev. Immunol., 12:775-808 (1994); Gerard, C. and Gerard, N. P., Curr. Opin. Immunol., 6:140-145
10 (1994)). The first receptor for the C-C chemokines that was cloned and expressed, binds the chemokines MIP-1 α and RANTES. Accordingly, this MIP-1 α /RANTES receptor was designated C-C chemokine receptor 1 (also referred to as CCR-1; Neote, K., et al., Cell, 72:415-425 (1993); Horuk, R. et al., WO 94/11504, May 26,
15 1994; Gao, J.-I. et al., J. Exp. Med., 177:1421-1427 (1993)). Two other receptors have been characterized which bind and/or signal in response to RANTES: CCR3 mediates binding and signaling of chemokines including eotaxin, RANTES, and MCP-3 (Ponath et al., J. Exp. Med., 183:2437 (1996)), and CCR5 binds chemokines including MIP-1 α , RANTES, MIP-1 β , and MCP-2. (Samson, et al., Biochem. 35: 3362-3367 (1996)).

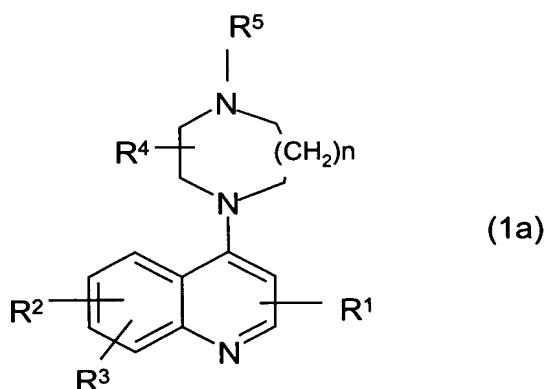
The CCR5 ligand RANTES is chemotactic for a variety of cell types, including
20 monocytes, eosinophils, and a subset of T-cells. The ability of RANTES to induce the directed migration of monocytes and a memory population of circulating T-cells (Schall, T. et al., Nature, 347:669-71 (1990)) suggests this chemokine and its receptor(s) plays an important role in chronic inflammatory diseases, since these diseases are characterized by destructive infiltrates of T cells and monocytes.
25 Macrophages and microglia also express CCR5 and the expression is upregulated following activation, especially under conditions of tissue damage such as demyelination. Particular diseases or conditions in which RANTES/CCR5 have been implicated include: transplant rejection, multiple sclerosis, arteriosclerosis (the formation of atherosclerotic plaques that lead to heart attacks and stroke), arthritis
30 (particularly osteo- and rheumatoid arthritis), stroke, atopic dermatitis, airway inflammatory disorders such as Rous Sarcoma Virus-induced bronchiolitis, delayed type hypersensitivity (DTH) reactions, glomerular nephritis, asthma, endometriosis

(Am J Obstet Gynecol 169(6): 1545-1549), cancers such as breast, cervical, melanoma, lymphoma, myeloma, Hodgkin's disease, and Hairy cell leukemia. (Cancer Research 62; 1093-1102)(Cancer Research 59; 4681-4687)(Clin Can Res. 7:285-289). Small molecule antagonists of CCR5 would have potential therapeutic utility in treating a wide range of disease states.

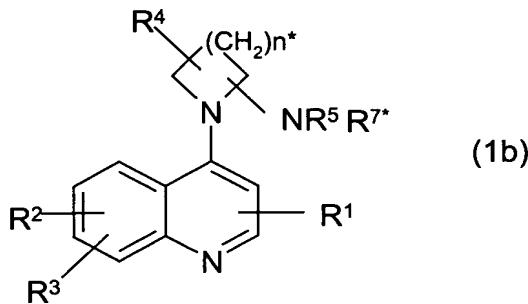
It has recently been recognized that the human immunodeficiency viruses (HIV) requires a chemokine receptor, most probably CCR5 or CXCR4, as well as the primary receptor CD4 for efficient entry into target cells (Levy, N. Engl. J. Med., 335(20), 1528-1530 (Nov. 14, 1996). The principal cofactor for entry mediated by the envelope glycoproteins of macrophage tropic strains of HIV-1 is CCR-5, a receptor for the chemokines RANTES, MIP-1 α , MIP-1 β , and MCP-2 (Deng, et al., Nature, 381, 661-666 (1996)). Accordingly, agents which could antagonize the interaction of HIV with chemokine receptors in humans should prevent infection in healthy individuals and slow or halt viral progression in infected patients leading to a viable method for the prevention or treatment of infection by HIV and the prevention or treatment of AIDS and AIDS related syndromes such as HIV dementia.

SUMMARY OF THE INVENTION

The present invention relates to CCR5 receptor antagonists of formulae (1a) or (1b):



or



enantiomers, diastereomers, salts and solvates thereof
wherein

R^1 and R^{1*} are independently hydrogen, a substituted or unsubstituted amino, alkyl,

5 haloalkyl, hydroxy, alkoxy or $-C(O)OR^{9a}$;

R^2 , R^{2*} , R^3 and R^{3*} are independently hydrogen, alkyl, halogenated alkyl, halogen, substituted or unsubstituted amino, nitro, cyano, or alkoxy;

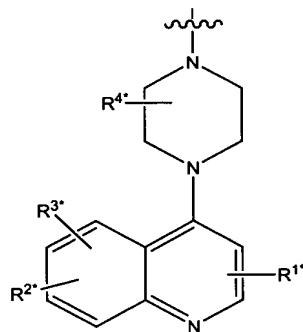
R^4 and R^{4*} are independently hydrogen or one or more alkyl groups;

R^5 is

10 (1) hydrogen, or

(2) R^9 , R^9 -aminocycloalkyl, R^9 -aminocycloalkenyl, (alkoxy)carbonyl, (aryloxy)carbonyl, $-SO_2R^9$, $-C(=O)-NR^7R^9$, $-C(=O)-NR^7-SO_2R^9$, $-C(=O)-R^6$, $-C(=O)-R^9$, $-C(=NR^{10})-R^9$, $-C(=S)-R^9$, $-C(=NR^{10})-NHR^9$, $-C(=S)-NHR^9$, or $-C(=S)-NR^7-SO_2R^9$ any of which can be substituted or unsubstituted;

15 R^6 is a group of formula



R^7 and R^{7*} are independently hydrogen, substituted or unsubstituted C₁₋₆ alkyl, or substituted or unsubstituted aryl;

R^9 is arylalkyl, cycloalkyl, cycloalkenyl, cycloalkylalkyl, alkyl, heterocyclalkyl, aryl or

20 heterocyclyl any of which can be substituted or unsubstituted;

R^{9a} is

(1) hydrogen, or

(2) arylalkyl, cycloalkyl, cycloalkenyl, cycloalkylalkyl, alkyl, heterocyclalkyl, aryl or heterocycl any of which can be substituted or unsubstituted;

R¹⁰ is

(1) hydrogen, or cyano;

5 (2) alkyl, or alkoxy, either of which optionally can be substituted;

n is 0,1,2 or 3; and

n* is 1, 2 or 3.

10 The compounds of the present invention are useful in the prevention and treatment of a wide variety of inflammatory and immunoregulatory disorders and diseases, allergic conditions, atopic conditions, as well as autoimmune, immunodeficiency, and demyelinating pathologies.

15 Also included in the invention are methods of using the compounds as agents for the treatment of CCR5 mediated disease states, in particular for the treatment of inflammatory diseases or conditions, autoimmune disorders, demyelinating pathologies, and immune deficiency disorders such as HIV infection.

20 In another aspect, the instant invention may be used to evaluate specific antagonists of CCR-5 receptors. Accordingly, the present invention is directed to the use of these compounds in the preparation and execution of screening assays for compounds which modulate the activity of CCR-5 receptors. For example, the compounds of this invention are useful for isolating receptor mutants, which are 25 excellent screening tools for more potent compounds. Furthermore, the compounds of this invention are useful in establishing or determining the binding site of other compounds to CCR-5 receptors, e.g., by competitive inhibition.

The compounds of the invention can be used in the treatment of mammals, 30 preferably humans, comprising administering to such mammal in need thereof, an effective amount of a compound of formula (I), or a pharmaceutically acceptable salt thereof, optionally in the form of a diastereomer and enantiomer.

Preferred compounds within the scope of the invention include compounds

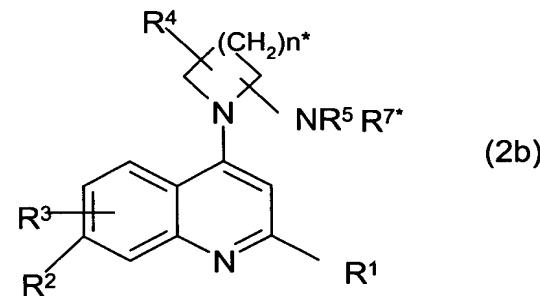
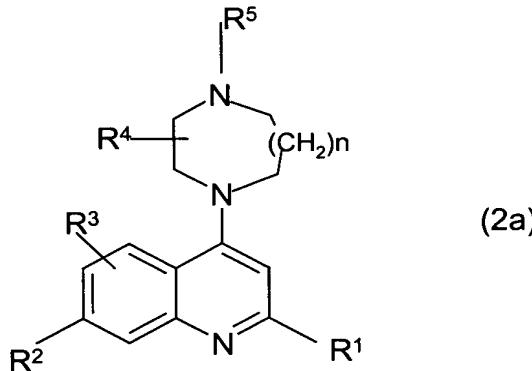
where

R^5 is alkoxycarbonyl (e.g., ethoxycarbonyl), $-C(=O)NHR^9$, $-C(=O)-NR^7-SO_2R^9$, $-C(=S)NHR^9$, or $-C(=S)-NR^7-SO_2R^9$; and

R^7 is hydrogen, substituted or unsubstituted C_{1-6} alkyl, or substituted or unsubstituted 5 aryl;

R^9 is alkyl, cycloalkyl, cycloalkylalkyl, aryl, arylalkyl, heterocyclyl, or heterocyclylalkyl any of which may be optionally substituted (especially preferred substituents are independently selected from alkyl and halo groups).

10 Preferred compounds within the scope of formulae (1a) and (1b) include compounds of the following formulae (2a) and (2b)



enantiomers, diastereomers, salts and solvates thereof

wherein

15 R^1 is hydrogen, amino or substituted amino;

R^2 is halo (especially chloro);

R^3 is hydrogen, alkyl, halogenated alkyl, halogen, substituted or unsubstituted amino, nitro, cyano, or alkoxy;

R^4 is hydrogen or one or more alkyl groups;

20 R^5 is alkoxycarbonyl, $-C(=O)NHR^9$, $-C(=O)-NR^7-SO_2R^9$, $-C(=S)NHR^9$ or $-C(=S)-NR^7-SO_2R^9$;

R^9 is cycloalkyl, cycloalkenyl, heterocyclyl, heterocyclylalkyl, aryl, or arylalkyl any of which may be optionally substituted.

25 Preferred compounds within the scope of formulae (2a) and (2b) include compounds wherein

R¹ is hydrogen or amino;

R² is halo;

R³ is hydrogen;

R⁴ is hydrogen;

5 R⁵ is -C(=O)NHR⁹, or -C(=S)NHR⁹;

R⁹ is heterocyclo (especially pyrrolidinyl, hexahydrothiazepinyl, tetrahydrobenzazepinyl, or hexahydroazepinyl) optionally substituted with one to three hydroxy, oxo or thioxo groups and further optionally substituted with one or more

10 (i) -(CR²⁰R²¹)_m-C(=O)R¹⁵, -(CR²⁰R²¹)_m-C(=O)OR¹⁵,
 -(CR²⁰R²¹)_m-SO₂R^{15a}, -(CR²⁰R²¹)_m-C(=O)NR¹⁶R¹⁷,
 -(CR²⁰R²¹)_m-C(=S)NR¹⁶R¹⁷,
 -(CR²⁰R²¹)_m-C(=O)NR¹⁶-SO₂R^{15b};

15 (ii) aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, cycloalkyl or cycloalkylalkyl any of which may be optionally independently substituted with one or more halo, alkoxy, hydroxy, or haloalkyl;

(iii) cyano;

R¹⁵ is hydrogen, alkyl, haloalkyl, alkenyl, cycloalkyl, cycloalkylalkyl, aryl, arylalkyl, aryloxyalkyl;

20 R^{15a} is hydrogen, alkyl, haloalkyl, alkenyl, cycloalkyl, cycloalkylalkyl, aryl, arylalkyl, aryloxyalkyl provided that when m is zero R^{15a} is not hydrogen;

R^{15b} is alkyl, haloalkyl, alkenyl, cycloalkyl, cycloalkylalkyl, aryl, arylalkyl, aryloxyalkyl;

25 R¹⁶ and R¹⁷ are independently hydrogen, alkyl, alkenyl, cycloalkyl, haloalkyl, hydroxyalkyl, alkoxyalkyl, aryloxyalkyl, alkenyl, cycloalkylalkyl, heterocyclyl, heterocyclylalkyl, aryl, arylalkyl;

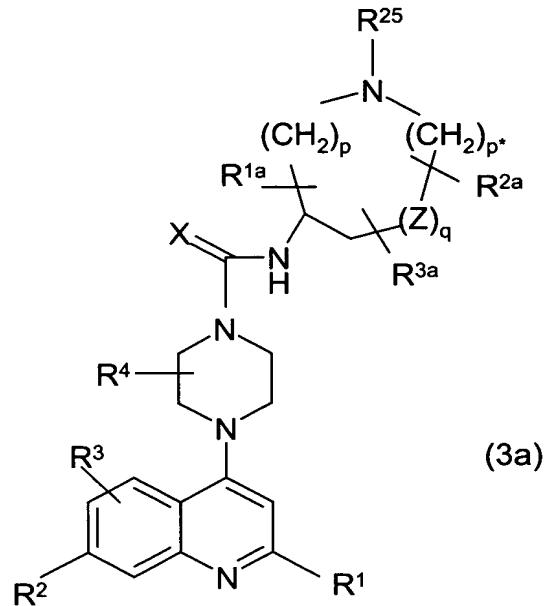
or R¹⁶ and R¹⁷ together with the nitrogen atom to which they are bonded may combine to form a heterocyclyl ring;

R²⁰ and R²¹ at each occurrence are the same or different and are independently selected from hydrogen or alkyl;

30 m is 0, 1, 2 or 3; and
 n is zero.

Preferred compounds within the scope of formula (2a) include compounds of

formula (3a)



wherein

 R^1 is hydrogen or amino;5 R^2 is halo; R^3 is hydrogen, alkyl, halogenated alkyl, halogen, substituted or unsubstituted amino, nitro, cyano, or alkoxy; R^4 is hydrogen or one or more alkyl groups; R^{1a} , R^{2a} and R^{3a} are independently selected from hydrogen, oxo, thioxo10 or when bonded to adjacent ring carbon atoms R^{2a} and R^{3a} may combine to form a fused aryl or heterocyclo ring; R^{25} is

(i) hydrogen, or cyano

(ii) alkyl, cycloalkyl, haloalkyl, hydroxyalkyl, alkoxyalkyl, aryloxyalkyl, alkenyl, cycloalkylalkyl, heterocyclalkyl, aryl, or arylalkyl any of which may be 15 optionally independently substituted with one to three halo, alkoxy, hydroxy, alkyl, or haloalkyl; or

(iii) $-(CR^{20}R^{21})_m-C(=O)R^{15}$, $-(CR^{20}R^{21})_m-C(=O)OR^{15}$, $-(CR^{20}R^{21})_m-SO_2R^{15a}$,
 $-(CR^{20}R^{21})_m-C(=O)NR^{16}R^{17}$, $-(CR^{20}R^{21})_m-C(=S)NR^{16}R^{17}$; or
 $-(CR^{20}R^{21})_m-C(=O)NR^{16}-SO_2R^{15b}$;20 R^{15} is hydrogen, alkyl, haloalkyl, alkenyl, cycloalkyl, cycloalkylalkyl, aryl, arylalkyl, aryloxyalkyl;

R^{15a} is hydrogen, alkyl, haloalkyl, alkenyl, cycloalkyl, cycloalkylalkyl, aryl, arylalkyl, aryloxyalkyl provided that when m is zero R^{15a} is not hydrogen;

R^{15b} is alkyl, haloalkyl, alkenyl, cycloalkyl, cycloalkylalkyl, aryl, arylalkyl, aryloxyalkyl;

R^{16} and R^{17} are independently hydrogen, alkyl, cycloalkyl, haloalkyl, hydroxyalkyl,

5 alkoxyalkyl, aryloxyalkyl, alkenyl, cycloalkylalkyl, heterocyclalkyl, aryl, or arylalkyl;

or R^{16} and R^{17} together with the nitrogen atom to which they are bonded may combine to form a heterocycl ring;

R^{20} and R^{21} at each occurrence are the same or different and are independently

10 selected from hydrogen or alkyl;

Z is $-S-$, $-S(O)-$ or $-S(O)_2-$;

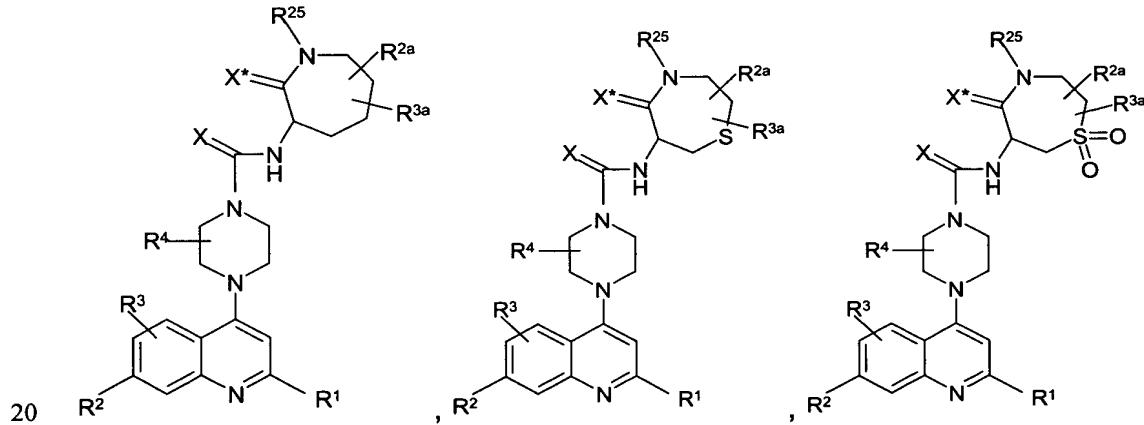
p is 1 or 2;

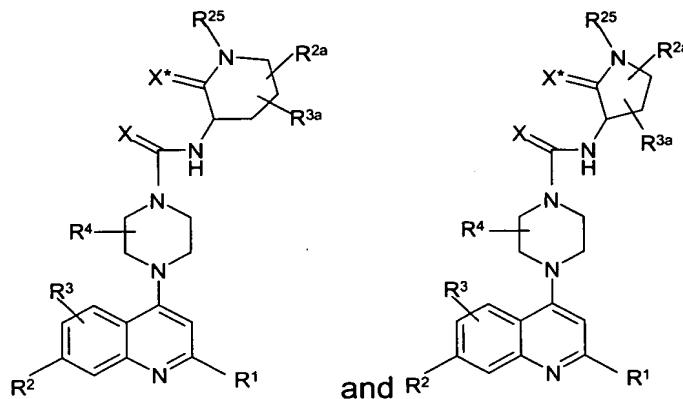
p^* is 0, 1, 2, 3 or 4;

q is 0 or 1;

15 m is 0, 1 or 2.

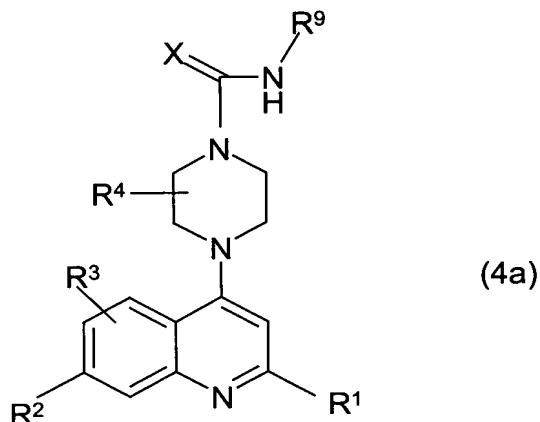
Preferred compounds within the scope of formula (3a) include compounds of the following formulae:





where X and X^* are independently O or S .

Preferred compounds within the scope of formula (2a) include compounds of
5 formula (4a)



wherein

R^1 is hydrogen or amino;

R^2 is halo;

10 R^3 is hydrogen, alkyl, halogenated alkyl, halogen, substituted or unsubstituted amino, nitro, cyano, or alkoxy;

R^4 is hydrogen or one or more alkyl groups;

R^9 is

(a) alkyl, haloalkyl, alkoxycarbonylalkyl, $-\text{CH}_2\text{C}(=\text{O})\text{OR}^{15}$, $-\text{CH}_2\text{C}(=\text{O})\text{R}^{15}$, or $-\text{CH}_2\text{C}(=\text{O})\text{NR}^{16}\text{R}^{17}$;

15 (b) cycloalkyl (especially cyclohexyl, cycloheptyl, benzofused-cyclohexyl, or benzofused-cycloheptyl) optionally substituted with one to three groups independently selected from alkyl, haloalkyl, alkoxy, aryloxy, arylalkyloxy,

heteroaryloxy, halo, hydroxy, oxo, thioxo, =N-OR¹⁵, -(CR²⁰R²¹)_m-C(=O)OR¹⁵, -(CR²⁰R²¹)_m-C(=O)R¹⁵, -(CR²⁰R²¹)_m-C(=O)NR¹⁶R¹⁷;

(c) aryl (especially phenyl or benzodioxinyl) optionally substituted with one to three groups independently selected from alkyl, alkoxy, heteroaryloxy, halo, 5 hydroxy, oxo, thioxo, =N-OR¹⁵, -(CR²⁰R²¹)_m-C(=O)OR¹⁵, -(CR²⁰R²¹)_m-C(=O)R¹⁵, -(CR²⁰R²¹)_m-C(=O)NR¹⁶R¹⁷; or

(d) heteroaryl (especially pyridyl) optionally substituted with one to three groups independently selected from alkyl, alkoxy, aryloxy, heteroaryloxy, halo, 10 hydroxy, oxo, thioxo, =N-OR¹⁵, -(CR²⁰R²¹)_m-C(=O)OR¹⁵, -(CR²⁰R²¹)_m-C(=O)R¹⁵, -(CR²⁰R²¹)_m-C(=O)NR¹⁶R¹⁷.

Preferred compounds within the scope of formulae (1a) and (1b) include those wherein R⁵ is -C(O)NHR⁹ wherein R⁹ is 2-oxopyrrolidinyl, 5-oxohexahydro-1,4-thiazepin-6-yl, 5-oxohexahydro-1,1-dioxido-1,4-thiazepin-6-yl, (2,3,4,5)-tetrahydro-2-15 oxo-1*H*-1-benzazepin-3-yl, hexahydro-1*H*-azepin-3-yl, 2-oxohexahydro-1*H*-azepin-3-yl, any of which may be optionally independently substituted by

- a) substituted or un-substituted alkyl, allyl, cyclopropylmethyl,
- b) substituted phenylmethyl;
- c) 2, 3, or 4-pyridinylmethyl
- 20 d) substituted phenyl
- e) cyano;
- f) formyl, alkycarbonyl;
- g) cycloalkylcarbonyl, heterocyclocarbonyl (preferably pyrrolidinylcarbonyl), or arylcarbonyl;
- 25 h) alkylsulfonyl, arylsulfonyl;
- i) alkoxy carbonyl, fluorinated alkoxy carbonyl, alloxy carbonyl;
- j) substituted phenoxy carbonyl, benzoxy carbonyl;
- k) aminocarbonyl, substituted or unsubstituted alkylaminocarbonyl, fluorinated 30 alkylaminocarbonyl, allylaminocarbonyl, cycloalkylaminocarbonyl;
- l) *N,N*-alkylalkylaminocarbonyl;
- m) alkylaminocarbonothionyl;
- n) substituted or unsubstituted phenylaminocarbonyl;
- o) (arylsulfonyl)aminocarbonyl;

Examples of specific preferred R⁵ groups include:

- 4-fluorophenylaminocarbonyl;
- 4-tert-butoxycarbonyl;
- 5 3,4-difluorophenylaminocarbonyl;
- 2,3,4-trifluorophenylaminocarbonyl;
- 4-phenylaminocarbonyl;
- 4-methylphenylaminocarbonyl;
- 3-methylphenylaminocarbonyl;
- 10 2-methylphenylaminocarbonyl;
(4-n-butylphenylaminocarbonyl) ;
- 4-methoxyphenylaminocarbonyl;
- 4-benzyloxyphenylaminocarbonyl;
- benzylaminocarbonyl;
- 15 4-phenethylaminocarbonyl;
phenylpropylaminocarbonyl;
(1-methoxycarbonyl-1-phenylethyl)aminocarbonyl;
- 1-phenylethylaminocarbonyl;
- 2-chlorobenzylaminocarbonyl;
- 20 2-methylbenzylaminocarbonyl;
2-trifluoromethylbenzylaminocarbonyl;
- 1-indanylaminocarbonyl;
- 1,2,3,4-tetrahydronaphth-1-ylaminocarbonyl;
- 4-pyridinylaminocarbonyl;
- 25 3-pyridinylaminocarbonyl;
- 2-pyridinylaminocarbonyl;
- 4-trifluoromethylphenylaminocarbonyl;
- 4-ethoxyphenylaminocarbonyl;
- 4-isopropylphenylaminocarbonyl;
- 30 4-acetylphenylaminocarbonyl;
- 4-dimethylaminophenylaminocarbonyl;
- cyclopentylaminocarbonyl;
- 3-piperonylaminocarbonyl;

(6-methylbenzothiazol-2-yl)phenylaminocarbonyl ;
cyclobutylaminocarbonyl;
cyclohexylaminocarbonyl;
adamant-1-ylaminocarbonyl;

5 exo-norborn-2-ylaminocarbonyl;
benzimidazol-2-yl;
pyrimidin-2-yl;
cycloheptylaminocarbonyl;
cyclooctylaminocarbonyl;

10 (4-methylcyclohexyl)aminocarbonyl;
(2-methylcyclohexyl)aminocarbonyl;
fluorophenylaminocarbonyl;
2,3-dimethylcyclohexylaminocarbonyl;
4-(trifluoromethyl)cyclohexylaminocarbonyl;

15 *cis*-4-methylcyclohexylaminocarbonyl;
trans-4-methylcyclohexylaminocarbonyl
3,4-dimethoxyphenylaminocarbonyl;
2,3-dihydro-1, 4-benzodioxin-6-ylaminocarbonyl;
4-fluoro-3-methoxyphenylaminocarbonyl;

20 3-fluoro-4-methoxyphenylaminocarbonyl;
3-chloro-4-methoxyphenylaminocarbonyl;
2-hydroxyphenylaminocarbonyl;
3-cyclohexen-1-ylaminocarbonyl;
3-fluoro-6,7,8,9-tetrahydro-5*H*-benzo[*α*]cyclohepten-5-ylaminocarbonyl;

25 6,7,8,9-tetrahydro-5*H*-benzo[*α*]cyclohepten-5-ylaminocarbonyl;
2-fluoro-6,7,8,9-tetrahydro-5*H*-benzo[*α*]cyclohepten-7-ylaminocarbonyl;
9-(acetoxy)-2-fluoro-6,7,8,9-tetrahydro-5*H*-benzo[*α*]cyclohepten-7-yl-aminocarbonyl;
2-fluoro-6,7,8,9-tetrahydro-9-hydroxy-5*H*-benzo[*α*]cyclohepten-7-yl-aminocarbonyl;

30 4-*tert*-butoxycarbonyl-hexahydro-1*H*-azepin-4-ylaminocarbonyl;
4-benzoxycarbonyl-hexahydro-1*H*-azepin-4-ylaminocarbonyl;
3-hydroxycycloheptylaminocarbonyl;

3-oxocycloheptylaminocarbonyl;
3-hydroxy-3-methylcycloheptylaminocarbonyl;
3-(ethoxyimino)cycloheptylaminocarbonyl;
3-methoxycycloheptylaminocarbonyl;

5 3-[(4-fluorophenyl)methoxy]cycloheptylaminocarbonyl;
2-pyridinyloxy)cycloheptylaminocarbonyl;
3-(2-pyrimidinyloxy)cycloheptylaminocarbonyl;
3-[(3-methyl-2-pyridinyl)oxy]cycloheptylaminocarbonyl;

10 3-[(4-methyl-2-pyridinyl)oxy]cycloheptylaminocarbonyl;
(3*R*)-hexahydro-2-oxo-1*H*-azepin-3-ylaminocarbonyl;
(3*S*)-hexahydro-2-oxo-1*H*-azepin-3-ylaminocarbonyl;

15 (3*S*)-hexahydro-2-oxo-1-methyl-1*H*-azepin-3-ylaminocarbonyl;
(3*S*)-hexahydro-2-oxo-1-(phenylmethyl)-1*H*-azepin-3-ylaminocarbonyl;
(3*S*)-hexahydro-2-oxo-1-[(4-fluorophenyl)methyl]-1*H*-azepin-3-ylaminocarbonyl;

20 (3*S*)-hexahydro-2-oxo-1-ethyl-1*H*-azepin-3-ylaminocarbonyl;
(3*S*)-hexahydro-2-oxo-1-(2-aminocarbonyl)ethyl]-1*H*-azepin-3-ylaminocarbonyl;
(3*S*)-hexahydro-2-oxo-1-(3-pyridinylmethyl)-1*H*-azepin-3-ylaminocarbonyl;

25 (3*S*)-hexahydro-2-oxo-1-(2-propenyl)-1*H*-azepin-3-ylaminocarbonyl;
(3*S*)-hexahydro-2-oxo-1-(cyclopropylmethyl)-1*H*-azepin-3-ylaminocarbonyl;
(3*S*)-hexahydro-2-oxo-1-(2-ethoxycarbonylethyl)-1*H*-azepin-3-ylaminocarbonyl;

30 (3*S*)-hexahydro-2-oxo-1-[(4-methoxyphenyl)methyl]-1*H*-azepin-3-ylaminocarbonyl;
4-fluorophenylmethylcarbonyl;
2,3,4,5,6-pentafluorophenylaminocarbonyl;
2-methoxyphenylaminocarbonyl;

25 1,2-dihydro-2-oxo-3-pyridinylaminocarbonyl;
cyclohexylaminocarbonothioyl;
2,2,2-trifluoroethylaminocarbonothioyl;
(2-ethoxycarbonyl)ethylaminocarbonothioyl;
(1*S*)-2-methoxy-1-methyl-2-oxoethylaminocarbonyl;

30 (1*S*)-2-(4-fluorophenylamino)-1-methyl-2-oxoethylaminocarbonyl;
2-[N-methyl(phenylmethyl)amino]-2-oxoethylaminocarbonyl;
2-[(4-fluorophenyl)amino]-2-oxoethylaminocarbonyl;
2-(dimethylamino)-2-oxoethylaminocarbonyl;

2-(methylphenylamino)-2-oxoethylaminocarbonyl;
4-nitrobenzoxycarbonyl;
2-[(methoxycarbonyl)cycloheptyl]imino]oxycarbonyl;
2-(methoxycarbonyl)cycloheptylaminocarbonyl;

5 (6R)-hexahydro-5-oxo-1,4-thiazepin-6-ylaminocarbonothioyl;
(6R)-hexahydro-5-oxo-1,4-thiazepin-6-ylaminocarbonyl;
(6R)-hexahydro-1,1-dioxido-5-oxo-1,4-thiazepin-6-ylaminocarbonyl;
(3S)-2-oxopyrrolidinylaminocarbonyl;
(3S)-2,3,4,5-tetrahydro-2-oxo-1*H*-1-benzazepin-3-ylaminocarbonyl;

10 (3S)-2,3,4,5-tetrahydro-1-methyl-2-oxo-1*H*-1-benzazepin-3-ylaminocarbonyl;
(3S)-1,2,3,4-tetrahydro-2-oxoquinolinylaminocarbonyl;
2-methoxy-3-pyridinylaminocarbonyl;
1,2-dihydro-1-methyl-2-oxo-3-pyridinylaminocarbonyl;
(3*R*)-hexahydro-2-thioxo-1*H*-azepin-3-ylaminocarbonyl;

15 (9S)-6,7,8,9-tetrahydro-5*H*-[1,2,4]triazolo[4,3-*α*]azepin-9-ylaminocarbonyl;
(9S)-2,5,6,7,8,9-hexahydro-3-oxo-3*H*-[1,2,4]triazolo[4,3-*α*]azepin-9-ylaminocarbonyl;
N-[(3S)-1-acetylhexahydro-1*H*-azepin-3-ylaminocarbonyl;
2-(methoxycarbonyl)cyclohexylaminocarbonyl;
2-(hydroxycarbonyl)cyclohexylaminocarbonyl;

20 2-[(methylamino)carbonyl]cyclohexylaminocarbonyl;
4-(2-amino-7-chloro-4-quinolinyl)-N-[(3S)-hexahydro-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide;
(3S)-1-[(2,6-dimethylphenyl)methyl]hexahydro-2-oxo-1*H*-azepin-3-ylaminocarbonyl;
(3S)-hexahydro-2-oxo-1-(2-pyridinylmethyl)-1*H*-azepin-3-ylaminocarbonyl;

25 (3S)-hexahydro-2-oxo-1-(4-pyridinylmethyl)-1*H*-azepin-3-ylaminocarbonyl;
(3S)-hexahydro-2-oxo-1-(2-hydroxy-3-phenoxypropyl)-1*H*-azepin-3-ylaminocarbonyl;
(3S)-hexahydro-1-(2-hydroxypropyl)-2-oxo-1*H*-azepin-3-ylaminocarbonyl;
(3S)-hexahydro-2-oxo-1-(2-methoxy-2-oxo-ethyl)-1*H*-azepin-3-ylaminocarbonyl;

30 (3S)-hexahydro-2-oxo-1-(2-hydroxy-2-oxo-ethyl)-1*H*-azepin-3-ylaminocarbonyl;
(3S)-hexahydro-2-oxo-1-(2-benzylamino-2-oxo-ethyl)-1*H*-azepin-3-ylaminocarbonyl;
(3S)-hexahydro-2-oxo-1-(2-methylamino-2-oxo-ethyl)-1*H*-azepin-3-ylaminocarbonyl;
(3S)-hexahydro-2-oxo-1-[2-(4-pyridinyl)amino-2-oxo-ethyl]-1*H*-azepin-3-ylaminocarbonyl;

(3*R*)-1-hexahydro-2-oxo-(4-fluorophenyl)-1*H*-azepin-3-ylaminocarbonyl;

(3*S*)-hexahydro-2-oxo-1-(2-phenoxyacetyl)-1*H*-azepin-3-ylaminocarbonyl;

(3*S*)-hexahydro-2-oxo-1-acetyl-1*H*-azepin-3-ylaminocarbonyl;

(3*S*)-hexahydro-2-oxo-1-cyclopropylcarbonyl-1*H*-azepin-3-ylaminocarbonyl;

5 (3*S*)-hexahydro-2-oxo-1-benzoyl-1*H*-azepin-3-ylaminocarbonyl
(3*S*)-hexahydro-2-oxo-1-ethylsulfonyl-1*H*-azepin-3-ylaminocarbonyl;

(3*S*)-hexahydro-2-oxo-1-phenylsulfonyl-1*H*-azepin-3-ylaminocarbonyl;

(3*S*)-hexahydro-2-oxo-1-[(phenylsulfonyl)amino]carbonyl]-1*H*-azepin-3-ylaminocarbonyl;

10 (3*S*)-hexahydro-2-oxo-1-(2-phenylacetyl)-1*H*-azepin-3-ylaminocarbonyl;
4-(2-amino-7-chloro-4-quinolinyl)-1-piperazinylaminocarbonyl;

(3*S*)-hexahydro-2-oxo-1-formyl-1*H*-azepin-3-ylaminocarbonyl;

(3*S*)-hexahydro-2-oxo-1-(2,2-dimethyl-1-oxopropyl)-1*H*-azepin-3-ylaminocarbonyl;

(3*S*)-hexahydro-2-oxo-1-(methylsulfonyl)-1*H*-azepin-3-ylaminocarbonyl;

15 (3*S*)-hexahydro-2-oxo-1-(methoxycarbonyl)-1*H*-azepin-3-ylaminocarbonyl;
(3*S*)-hexahydro-2-oxo-1-(ethoxycarbonyl)-1*H*-azepin-3-ylaminocarbonyl;

(3*S*)-hexahydro-2-oxo-1-(2,2,2-trifluoroethoxycarbonyl)-1*H*-azepin-3-ylaminocarbonyl;

(3*S*)-hexahydro-2-oxo-1-(2-propenoxy carbonyl)-1*H*-azepin-3-ylaminocarbonyl;

(3*S*)-hexahydro-2-oxo-1-(phenoxy carbonyl)-1*H*-azepin-3-ylaminocarbonyl;

20 (3*S*)-hexahydro-2-oxo-1-(4-fluorophenoxy carbonyl)-1*H*-azepin-3-ylaminocarbonyl;
(3*S*)-hexahydro-2-oxo-1-(benzoxycarbonyl)-1*H*-azepin-3-ylaminocarbonyl;

(3*S*)-hexahydro-2-oxo-1-cyano-1*H*-azepin-3-ylaminocarbonyl;

(3*S*)-hexahydro-2-oxo-1-(phenoxy carbonyl)-1*H*-azepin-3-ylaminocarbonyl;

(3*S*)-hexahydro-2-oxo-1-(cyclohexylaminocarbonyl)-1*H*-azepin-3-ylaminocarbonyl;

25 (3*S*)-hexahydro-2-oxo-1-(aminocarbonyl)-1*H*-azepin-3-ylaminocarbonyl;
(2*S*)-6-[(trichloroacetyl)amino]carbonyl]amino]-1-methoxy-1-oxo-hexyl-2-aminocarbonyl;

(3*S*)-hexahydro-2-oxo-1-(2,2,2-trifluoroethylaminocarbonyl)-1*H*-azepin-3-ylaminocarbonyl;

30 (3*S*)-hexahydro-2-oxo-1-(cyclopropylaminocarbonyl)-1*H*-azepin-3-ylaminocarbonyl;
(3*S*)-hexahydro-2-oxo-1-(propylaminocarbonyl)-1*H*-azepin-3-ylaminocarbonyl;

(3*S*)-hexahydro-2-oxo-1-(pyrrolidinylcarbonyl)-1*H*-azepin-3-ylaminocarbonyl;

(3*S*)-hexahydro-2-oxo-1-(cyclopentylaminocarbonyl)-1*H*-azepin-3-ylaminocarbonyl;

(3S)-hexahydro-2-oxo-1-(cyclobutylaminocarbonyl)-1*H*-azepin-3-ylaminocarbonyl;
 (3S)-hexahydro-2-oxo-1-(4-fluorophenylaminocarbonyl)-1*H*-azepin-3-ylaminocarbonyl;
 (3S)-hexahydro-2-oxo-1-(phenylaminocarbonyl)-1*H*-azepin-3-ylaminocarbonyl;
 (3S)-hexahydro-2-oxo-1-(isopropylaminocarbonyl)-1*H*-azepin-3-ylaminocarbonyl;

5 (3S)-hexahydro-2-oxo-1-(n-butylaminocarbonyl)-1*H*-azepin-3-ylaminocarbonyl;
 (3S)-hexahydro-2-oxo-1-(2-methylpropylaminocarbonyl)-1*H*-azepin-3-ylaminocarbonyl;
 (3S)-hexahydro-2-oxo-1-(t-butylaminocarbonyl)-1*H*-azepin-3-ylaminocarbonyl;
 (3S)-hexahydro-2-oxo-1-(*N,N*-dimethylaminocarbonyl)-1*H*-azepin-3-ylaminocarbonyl;
 (3S)-hexahydro-2-oxo-1-(4-trifluoromethylaminocarbonyl)-1*H*-azepin-3-

10 ylaminocarbonyl;
 (3S)-hexahydro-2-oxo-1-(3-fluorophenylaminocarbonyl)-1*H*-azepin-3-ylaminocarbonyl;
 (3S)-hexahydro-2-oxo-1-(4-methylphenylaminocarbonyl)-1*H*-azepin-3-
 ylaminocarbonyl;
 (3S)-hexahydro-2-oxo-1-(2-fluorophenylaminocarbonyl)-1*H*-azepin-3-ylaminocarbonyl;

15 (3S)-hexahydro-2-oxo-1-(*N*-2,6-diethylphenylaminocarbonyl)-1*H*-azepin-3-
 ylaminocarbonyl;
 (3S)-hexahydro-2-oxo-1-(*N*-methylaminocarbonothioyl)-1*H*-azepin-3-ylaminocarbonyl;
 (3S)-hexahydro-2-oxo-1-(*N*-ethylaminocarbonothioyl)-1*H*-azepin-3-ylaminocarbonyl;
 (3S)-hexahydro-2-oxo-1-(*N*-isopropylaminocarbonothioyl)-1*H*-azepin-3-

20 ylaminocarbonyl;
 (3S)-hexahydro-2-oxo-1-(*N*-2-propenylaminocarbonothioyl)-1*H*-azepin-3-
 ylaminocarbonyl;
 (3S)-hexahydro-2-oxo-1-(*N*-methylaminocarbonyl)-1*H*-azepin-3-ylaminocarbonyl;
 (3S)-hexahydro-2-oxo-1-[*N*-(2-propenyl)aminocarbonyl]-1*H*-azepin-3-ylaminocarbonyl;

25 (3S)-hexahydro-2-oxo-1-[*N*-(2-chloroethyl)aminocarbonyl]-1*H*-azepin-3-
 ylaminocarbonyl;
 (3R)-hexahydro-2-oxo-1-(*N*-methylaminocarbonyl)-1*H*-azepin-3-ylaminocarbonyl;

Also preferred are compounds of formula 1a wherein R⁵ is

30 fluorophenylaminocarbonothionyl,
 cyanimino(phenoxy)methyl,
 cyanimino(ethoxy)methyl,
 cyanimino(4-fluorophenylamino)methyl,

cyanimino(4-fluorobenzylamino)methyl,
tert-butylamino(cyanimino)methyl,
cyanimino(ethylamino)methyl,
4-fluorophenylamino-2,3-dioxocyclobut-1-enyl,
5 ethoxyimino(4-fluorophenylamino)methyl,
methoxyimino(4-fluorophenylamino)methyl;
fluorobenzylamino(imino)methyl, or
fluorophenylamino(imino)methyl.

10 Also preferred are compounds wherein R¹ is

- a) -CO₂H
- b) -CONMe₂,
- c) methyl,
- d) -COOethyl,
- e) -H,
- f) substituted or unsubstituted amino;
- g) substituted (phenylamino)carbonylamino; or
- h) [2-oxohexahydro-1-(t-butyloxycarbonyl)-1*H*-azepin-3-yl]carbonylamino;

20 Also preferred are compounds wherein R² and R³ may be the same or different and are independently selected from

- a)-Cl
- b)-F
- c)-Br

25 e)-CF₃

f)-OCH₃

g)-Me

h) R² and R³ are on adjacent carbon atoms and combine to form -OCH₂O-

i)-NO₂

30 j)-NH₂

k)-NHAc

l)-NHMe or

m)-CN

Other preferred embodiments of the present invention include:

- a) A pharmaceutical composition comprising one or more compounds of formulae (1a) and/or (1b) in admixture with a pharmaceutically acceptable excipient, diluent, or carrier.
- b) A method for modulation of chemokine receptor activity in a mammal which comprises the administration of an effective amount of at least one compound of formulae (1a) or (1b).
- c) A method for the prevention or treatment of an inflammatory and immunoregulatory disorder or disease which comprises the administration to a patient of an effective amount of at least one compound of formulae (1a) or (1b).
- d) A method for the prevention or treatment of asthma, allergic rhinitis, dermatitis, conjunctivitis, or atherosclerosis which comprises the administration to a patient an effective amount of at least one compound of formulae (1a) or (1b).
- e) A method for the prevention or treatment of rheumatoid arthritis which comprises the administration to a patient of an effective amount of at least one compound of formulae (1a) or (1b).
- f) A method for preventing infection by HIV, treating infection by HIV, delaying of the onset of AIDS, or treating AIDS or AIDS dementia complex comprising the administration to a patient of an effective amount of at least one compound of formulae (1a) or (1b).
- g) A method for the prevention or treatment of demyelinating disorders such as multiple sclerosis or optic neuritis which comprises the administration to a patient of an effective amount of at least one compound of formulae (1a) or (1b).
- h) A method for the prevention or treatment of endometriosis which comprises the

administration to a patient of an effective amount of at least one compound of formulae (1a) or (1b).

i) A method for the prevention or treatment of ischemic stroke which comprises the

5 administration to a patient of an effective amount of at least one compound of formulae (1a) or (1b).

j) A method for the prevention or treatment of cancer which comprises the

administration to a patient of an effective amount of at least one compound of formulae
10 (1a) or (1b).

k) A method for the prevention or treatment of psoriasis which comprises the

administration to a patient of an effective amount of at least one compound of formulae
15 (1a) or (1b).

Preferred compounds of the present invention include:

7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline;

6-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline;

7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-methylquinoline;

20 5-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-methylquinoline;

7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-8-methylquinoline;

7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-3-methylquinoline;

4-[4-(4-Fluorophenylaminocarbonyl)piperazin-1-yl]-7-methoxy-2-methylquinoline;

7-Chloro-6-fluoro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline;

25 7-Fluoro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline;

6,7-Difluoro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline;

7-Cyano-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline;

4-[4-(4-Fluorophenylaminocarbonyl)piperazin-1-yl]-6,7-methylenedioxyquinoline;

4-[4-(4-Fluorophenylaminocarbonyl)piperazin-1-yl]-7-methylquinoline trifluoroacetate;

30 4-[4-(tert-Butoxycarbonyl)piperazin-1-yl]-7-nitroquinoline;

4-[4-(4-Fluorophenylaminocarbonyl)piperazin-1-yl]-7-nitroquinoline trifluoroacetate;

7-Amino-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline;

7-Acetylamino-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline,

trifluoroacetate;

Ethyl 4-[4-(tert-butoxycarbonyl)piperazin-1-yl]-7-chloroquinoline-2-carboxylate;

Ethyl 7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline-2-carboxylate;

5 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-(hydroxymethyl)quinoline, trifluoroacetate;

2-Azidomethyl-7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline;

7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-(methoxymethyl)quinoline, trifluoroacetate;

10 2-Aminomethyl-7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline, trifluoroacetate;

7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline-2-carboxylic acid;

2-tert-Butoxycarbonylamino-7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline;

15 2-Amino-7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline, trifluoroacetate

2-Acetylamino-7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline;

N-[7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinolin-2-yl]urea;

N-[7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinolin-2-yl]-N'-phenylurea;

20 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-(methylamino)quinoline, trifluoroacetate;

4-[4-(tert-Butoxycarbonyl)piperazin-1-yl]-7-chloro-2-quinolone;

7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-quinolone;

25 7-Chloro-2-dimethylamino-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline;

7-Chloro-2-ethylamino-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline;

7-Chloro-2-[2-(dimethylamino)ethylamino]-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline;

7-Chloro-2-cyclohexylamino-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-

30 yl]quinoline;

7-Chloro-2-cyclopropylamino-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline;

7-Chloro-2-cyclopropylmethylamino-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-

yl]quinoline;
4-[4-(tert-Butoxycarbonyl)piperazin-1-yl]-7-chloro-2-(2-hydroxyethylamino)quinoline;
7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-[2-(4-fluorophenylaminocarbonyloxy)ethylamino]quinoline;

5 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-(2-hydroxyethylamino)quinoline trifluoroacetate;
7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-(2-methoxyethyl)aminoquinoline;
4-[4-(tert-Butoxycarbonyl)piperazin-1-yl]-7-chloro-2-(n-propylamino)quinoline;

10 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-(n-propylamino)quinoline;
4-[4-(tert-Butoxycarbonyl)piperazin-1-yl]-7-chloro-2-(isopropylamino)quinoline
7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-(isopropylamino)quinoline;

15 4-[4-(tert-Butoxycarbonyl)piperazin-1-yl]-2-(n-butylamino)-7-chloroquinoline;
2-n-Butylamino-7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline;
4-[4-(tert-Butoxycarbonyl)piperazin-1-yl]-7-chloro-2-(n-pentylamino)-quinoline;
7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-(n-pentylamino)quinoline;

20 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-[2-(4-fluorophenyl)ethylamino]quinoline;
7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)-3-methylpiperazin-1-yl]quinoline;
4-[4-(tert-Butoxycarbonyl)-2-methylpiperazin-1-yl]-7-chloroquinoline;
7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)-2-methylpiperazin-1-yl]quinoline;

25 7-Chloro-4-[3,5-dimethyl-4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline;
7-Chloro-4-[3,6-dimethyl-4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline, trifluoroacetate;
7-Chloro-4-[5-(4-fluorophenylaminocarbonyl)-2,5-diazanorbornan-2-yl]quinoline, trifluoroacetate;

30 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)-1,4-diazepin-1-yl]quinoline;
7-Chloro-4-[4-(3,4-difluorophenylaminocarbonyl)piperazin-1-yl]quinoline;
7-Chloro-4-[4-(2,3,4-trifluorophenylaminocarbonyl)piperazin-1-yl]quinoline;
7-Chloro-4-[phenylaminocarbonyl)piperazin-1-yl]quinoline;

7-Chloro-4-[4-(4-methylphenylaminocarbonyl)piperazin-1-yl]quinoline;
7-Chloro-4-[4-(3-methylphenylaminocarbonyl)piperazin-1-yl]quinoline;
7-Chloro-4-[4-(2-methylphenylaminocarbonyl)piperazin-1-yl]quinoline;
4-[4-(4-n-Butylphenylaminocarbonyl)piperazin-1-yl]-7-chloro-quinoline;
5 7-Chloro-4-[4-(4-methoxyphenylaminocarbonyl)piperazin-1-yl]quinoline;
4-[4-(4-Benzylaminocarbonyl)piperazin-1-yl]-7-chloroquinoline;
4-[4-(Benzylaminocarbonyl)piperazin-1-yl]-7-chloroquinoline;
7-Chloro-4-[4-(phenethylaminocarbonyl)piperazin-1-yl]quinoline;
7-Chloro-4-[4-(phenylpropylaminocarbonyl)piperazin-1-yl]quinoline;
10 7-Chloro-4-[4-[(1-methoxycarbonyl-1-phenylethyl)aminocarbonyl]piperazin-1-yl]quinoline;
7-Chloro-4-[4-(1-phenylethylaminocarbonyl)piperazin-1-yl]quinoline;
7-Chloro-4-[4-(2-chlorobenzylaminocarbonyl)piperazin-1-yl]quinoline;
7-Chloro-4-[4-(2-methylbenzylaminocarbonyl)piperazin-1-yl]quinoline;
15 7-Chloro-4-[4-(2-trifluoromethylbenzylaminocarbonyl)piperazin-1-yl]quinoline;
7-Chloro-4-[4-(1-indanylaminocarbonyl)piperazin-1-yl]quinoline;
7-Chloro-4-[4-(1,2,3,4-tetrahydronaphth-1-ylaminocarbonyl)piperazin-1-yl]quinoline;
7-Chloro-4-[4-(4-pyridinylaminocarbonyl)piperazin-1-yl]quinoline;
7-Chloro-4-[4-(3-pyridinylaminocarbonyl)piperazin-1-yl]quinoline;
20 7-Chloro-4-[4-(2-pyridinylaminocarbonyl)piperazin-1-yl]quinoline;
7-Chloro-4-[4-(4-trifluoromethylphenylaminocarbonyl)piperazin-1-yl]quinoline;
7-Chloro-4-[4-(4-ethoxyphenylaminocarbonyl)piperazin-1-yl]quinoline;
7-Chloro-4-[4-(4-isopropylphenylaminocarbonyl)piperazin-1-yl]quinoline;
4-[4-(4-Acetylphenylaminocarbonyl)piperazin-1-yl]-7-chloroquinoline;
25 7-Chloro-4-[4-(4-dimethylaminophenylaminocarbonyl)piperazin-1-yl]quinoline;
7-Chloro-4-[4-(cyclopentylaminocarbonyl)piperazin-1-yl]quinoline;
7-Chloro-4-[4-(3-piperonylaminocarbonyl)piperazin-1-yl]quinoline;
7-Chloro-4-[4-[4-(6-methylbenzothiazol-2-yl)phenylaminocarbonyl]piperazin-1-yl]quinoline;
30 7-Chloro-4-[4-(cyclobutylaminocarbonyl)piperazin-1-yl]quinoline;
7-Chloro-4-[4-(cyclohexylaminocarbonyl)piperazin-1-yl]quinoline;
4-[4-(Adamant-1-ylaminocarbonyl)piperazin-1-yl]-7-chloroquinoline;
7-Chloro-4-[4-(exo-norborn-2-ylaminocarbonyl)piperazin-1-yl]quinoline;

7-Chloro-4-[4-(4-fluorophenylsulfonyl)piperazin-1-yl]quinoline;
7-Chloro-4-[4-(4-fluorophenylaminothiocarbonyl)piperazin-1-yl]quinoline;
7-Chloro-4-[4-[cyanimino(phenoxy)methyl]piperazin-1-yl]quinoline;
7-Chloro-4-[4-[cyanimino(ethoxy)methyl]piperazin-1-yl]quinoline;
5 7-Chloro-4-[4-(cyanimino(4-fluorophenylamino)methyl)piperazin-1-yl]quinoline;
7-Chloro-4-[4-(cyanimino(4-fluorobenzylamino)methyl)piperazin-1-yl]quinoline;
4-[4-(tert-Butylamino(cyanimino)methyl)piperazin-1-yl]-7-chloroquinoline;
7-Chloro-4-[4-[cyanimino(ethylamino)methyl]piperazin-1-yl]quinoline;
7-Chloro-4-[4-[4-(4-fluorophenylamino)-2,3-dioxocyclobut-1-enyl]piperazin-1-
10 yl]quinoline;
4-[4-(Benzimidazol-2-yl)piperazin-1-yl]-7-chloroquinoline;
7-Chloro-4-[4-(pyrimidin-2-yl)piperazin-1-yl]-7-chloroquinoline;
7-Chloro-4-[4-[ethoxyimino(4-fluorophenylamino)methyl]piperazin-1-yl]quinoline;
7-Chloro-4-[4-[methoxyimino(4-fluorophenylamino)methyl]piperazin-1-yl]quinoline;
15 7-Chloro-4-(piperazin-1-yl)quinoline;
7-Chloro-4-[4-(4-fluorobenzylamino(imino)methyl)piperazin-1-yl]quinoline;
7-Chloro-4-[4-(4-fluorophenylamino(imino)methyl)piperazin-1-yl]quinoline;
7-Chloro-4-[4-(cycloheptylaminocarbonyl)piperazin-1-yl]quinoline;
7-Chloro-4-[4-(cyclooctylaminocarbonyl)piperazin-1-yl]quinoline;
20 7-Chloro-4-[4-[(4-methylcyclohexyl)aminocarbonyl]piperazin-1-yl]quinoline;
-7-Chloro-4-[4-[(2-methylcyclohexyl)aminocarbonyl]piperazin-1-yl]quinoline;
2-(2-Aminoethylamino)-7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-
yl]quinoline;
7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-[2-
25 (trifluoroacetamino)ethylamino]quinoline;
7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-methoxyquinoline;
4-(7-Chloro-4-quinoliny)-N-(2,3-dimethylcyclohexyl)-1-piperazinecarboxamide;
4-(7-Chloro-4-quinoliny)-N-[4-(trifluoromethyl)cyclohexyl]-1-piperazinecarboxamide;
4-(7-Chloro-4-quinoliny)-N-(cis-4-methylcyclohexyl)-1-piperazinecarboxamide;
30 4-(7-Chloro-4-quinoliny)-N-(trans-4-methylcyclohexyl)-1-piperazinecarboxamide
4-(7-Chloro-4-quinoliny)-N-(3,4-dimethoxyphenyl)-1-piperazinecarboxamide;
4-(7-Chloro-4-quinoliny)-N-(2,3-dihydro-1,4-benzodioxin-6-yl)-1-
piperazinecarboxamide;

4-(7-Chloro-4-quinolinyl)-*N*-(4-fluoro-3-methoxyphenyl)-1-piperazinecarboxamide;
4-(7-Chloro-4-quinolinyl)-*N*-(3-fluoro-4-methoxyphenyl)-1-piperazinecarboxamide;
4-(7-Chloro-4-quinolinyl)-*N*-(3-chloro-4-methoxyphenyl)-1-piperazinecarboxamide;
4-(7-Chloro-4-quinolinyl)-*N*-(2-hydroxyphenyl)-1-piperazinecarboxamide;

5 4-(7-Chloro-4-quinolinyl)-*N*-(3-cyclohexen-1-yl)-1-piperazinecarboxamide;
4-(7-Chloro-4-quinolinyl)-*N*-(3-fluoro-6,7,8,9-tetrahydro-5*H*-benzo[α]cyclohepten-5-yl)-1-piperazinecarboxamide;
4-(7-Chloro-4-quinolinyl)-*N*-(6,7,8,9-tetrahydro-5*H*-benzo[α]cyclohepten-5-yl)-1-piperazinecarboxamide;

10 4-(7-Chloro-4-quinolinyl)-*N*-(2-fluoro-6,7,8,9-tetrahydro-5*H*-benzo[α]cyclohepten-7-yl)-1-piperazinecarboxamide;
4-(7-Chloro-4-quinolinyl)-*N*-[9-(acetyloxy)-2-fluoro-6,7,8,9-tetrahydro-5*H*-benzo[α]cyclohepten-7-yl]-1-piperazinecarboxamide;
4-(7-Chloro-4-quinolinyl)-*N*-(2-fluoro-6,7,8,9-tetrahydro-9-hydroxy-5*H*-benzo[α]cyclohepten-7-yl)-1-piperazinecarboxamide;

15 4-[[[4-(7-Chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-1*H*-azepine-1-carboxylic acid, 1,1-dimethylethyl ester;
4-[[[4-(7-Chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-1*H*-azepine-1-carboxylic acid, methyl ester;

20 4-[[[4-(7-Chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-1*H*-azepine-1-carboxylic acid, phenylmethyl ester;
4-(7-Chloro-4-quinolinyl)-*N*-(3-hydroxycycloheptyl)-1-piperazinecarboxamide;
4-(7-Chloro-4-quinolinyl)-*N*-(3-oxocycloheptyl)-1-piperazinecarboxamide;

25 4-(7-Chloro-4-quinolinyl)-*N*-(3-hydroxy-3-methylcycloheptyl)-1-piperazinecarboxamide;
4-(7-Chloro-4-quinolinyl)-*N*-[(3*E*)-3-(ethoxyimino)cycloheptyl]-1-piperazinecarboxamide;
4-(7-Chloro-4-quinolinyl)-*N*-(3-methoxycycloheptyl)-1-piperazinecarboxamide;
4-(7-Chloro-4-quinolinyl)-*N*-[3-[(4-fluorophenyl)methoxy]cycloheptyl]-1-piperazinecarboxamide;

30 4-(7-Chloro-4-quinolinyl)-*N*-[3-(2-pyridinyloxy)cycloheptyl]-1-piperazinecarboxamide;
4-(7-Chloro-4-quinolinyl)-*N*-[3-(2-pyrimidinyloxy)cycloheptyl]-1-piperazinecarboxamide;

4-(7-Chloro-4-quinolinyl)-*N*-[3-[(3-methyl-2-pyridinyl)oxy]cycloheptyl]-1-piperazinecarboxamide;

4-(7-Chloro-4-quinolinyl)-*N*-[3-[(4-methyl-2-pyridinyl)oxy]cycloheptyl]-1-piperazinecarboxamide;

5 4-(7-Chloro-4-quinolinyl)-*N*-[(3*R*)-hexahydro-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

4-(7-Chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

4-(7-Chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-1-methyl-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

10 4-(7-Chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-2-oxo-1-(phenylmethyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

4-(7-Chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-2-oxo-1-[(4-fluorophenyl)methyl]-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

15 4-(7-Chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-2-oxo-1-ethyl-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

(3*S*)-[[4-(7-Chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-acetamide;

4-(7-Chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-2-oxo-1-(3-pyridinylmethyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

20 4-(7-Chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-2-oxo-1-(2-propenyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

4-(7-Chloro-4-quinolinyl)-*N*-[(3*S*)-1-hexahydro-2-oxo-1-(cyclopropylmethyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

25 4-(7-Chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-2-oxo-1-[(4-methoxyphenyl)methyl]-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

(3*S*)-[[4-(7-Chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-acetic acid, ethyl ester;

4-(7-Chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-2-oxo-1-[3-(4-morpholinyl)propyl]-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

30 7-Chloro-4-(1-piperazinyl)-2-quinolinamine;

7-Chloro-4-[4-[(4-fluorophenyl)acetyl]-1-piperazinyl]-2-quinolinamine;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-(2,3,4,5,6-pentafluorophenyl)-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-(2-methoxyphenyl)-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-ethyl-1-piperazinecarboxamide;

5 4-(2-Amino-7-chloro-4-quinolinyl)-*N*-(4-methylphenyl)-1-piperazinecarboxamide;

4-[7-Chloro-2-[[[[(4-methylphenyl)amino]carbonyl]amino]-4-quinolinyl]-*N*-(4-methylphenyl)-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-tricyclo[3.3.1.1^{3,7}]dec-2-yl-1-piperazinecarboxamide;

10 4-(2-Amino-7-chloro-4-quinolinyl)-*N*-(1,2-dihydro-2-oxo-3-pyridinyl)-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-cyclohexyl-1-piperazinecarbothioamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-(2,2,2-trifluoroethyl)-1-piperazinecarbothioamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-(4-fluorophenyl)-1-piperazinecarbothioamide;

15 [[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonothioyl]amino]-acetic acid, ethyl ester;

(2S)-2-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-propanoic acid, methyl ester;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(1*S*)-2-[(4-fluorophenyl)amino]-1-methyl-2-oxoethyl]-1-piperazinecarboxamide;

20 4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[2-[methyl(phenylmethyl)amino]-2-oxoethyl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[2-[(4-fluorophenyl)amino]-2-oxoethyl]-1-piperazinecarboxamide;

25 4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[2-(dimethylamino)-2-oxoethyl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[2-(methylphenylamino)-2-oxoethyl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinecarboxylic acid, 4-nitrophenyl ester;

30 2-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]oxy]imino]-cycloheptanecarboxylic acid, methyl ester;

2-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-cycloheptanecarboxylic acid, methyl ester;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(6*R*)-hexahydro-5-oxo-1,4-thiazepin-6-yl]-1-piperazinecarbothioamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(6*R*)-hexahydro-5-oxo-1,4-thiazepin-6-yl]-1-piperazinecarboxamide;

5 4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(6*R*)-hexahydro-1,1-dioxido-5-oxo-1,4-thiazepin-6-yl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(3*S*)-2-oxopyrrolidinyl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(3*S*)-2,3,4,5-tetrahydro-2-oxo-1*H*-1-benzazepin-3-yl]-1-piperazinecarboxamide;

10 4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(3*S*)-2,3,4,5-tetrahydro-1-methyl-2-oxo-1*H*-1-benzazepin-3-yl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(3*S*)-1,2,3,4-tetrahydro-2-oxoquinolinyl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-cycloheptyl-1-piperazinecarboxamide;

15 4-(2-Amino-7-chloro-4-quinolinyl)-*N*-(4-fluoro-3-methoxyphenyl)-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-(2-methoxy-3-pyridinyl)-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-(1,2-dihydro-1-methyl-2-oxo-3-pyridinyl)-1-piperazinecarboxamide;

20 4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(3*R*)-hexahydro-2-thioxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(9*S*)-6,7,8,9-tetrahydro-5*H*-[1,2,4]triazolo[4,3- α]azepin-9-yl]-1-piperazinecarboxamide

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(9*S*)-2,5,6,7,8,9-hexahydro-3-oxo-3*H*-[1,2,4]triazolo[4,3- α]azepin-9-yl]-1-piperazinecarboxamide

25 4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-1-acetyl-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-cyclohexanecarboxylic acid, methyl ester;

30 3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-cyclohexanecarboxylic acid;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[3-[(methylamino)carbonyl]cyclohexyl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-hexahydro-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-hexahydro-2-oxo-1-methyl-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

5 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-hexahydro-2-oxo-1-(cyclopropylmethyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-hexahydro-2-oxo-1-(phenylmethyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-hexahydro-2-oxo-1-[(2,6-dimethylphenyl)methyl]-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

10 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-hexahydro-2-oxo-1-(2-pyridinylmethyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-hexahydro-2-oxo-1-(3-pyridinylmethyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

15 4-(2-amino-7-chloro-4-quinolinyl)-N-[(3S)-hexahydro-2-oxo-1-(4-pyridinylmethyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-hexahydro-2-oxo-1-(2-hydroxy-3-phenoxypropyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-hexahydro-2-oxo-1-(2-hydroxypropyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

20 (3S)-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine, 1-acetic acid, methyl ester;

(3S)-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine, 1-acetic acid;

25 (3S)-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-N-(1-phenylmethyl)-1*H*-azepine-1-acetamide;

(3S)-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-N-methyl-2-oxo-1*H*-azepine-1-acetamide;

(3S)-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-N-(4-pyridinyl)-1*H*-azepine-1-acetamide;

30 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3*R*)-1-(4-fluorophenyl)hexahydro-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

4-[7-Chloro-2-(methylamino)-4-quinolinyl]-*N*-[(3*S*)-hexahydro-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-2-oxo-1-(2-phenoxyacetyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

5 4-(7-Chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-2-oxo-1-acetyl-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-2-oxo-1-acetyl-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

10 4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-2-oxo-1-(cyclopropylcarbonyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-2-oxo-1-benzoyl-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

15 4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-2-oxo-1-(ethylsulfonyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-2-oxo-1-(phenylsulfonyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

20 4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-2-oxo-1-[(phenylsulfonyl)amino]carbonyl]-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-2-oxo-1-(2-phenylacetyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

25 4-[4-[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]-1-piperazinyl]-7-chloro-2-quinolinamine;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-2-oxo-1-formyl-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

30 4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-2-oxo-1-(2,2-dimethyl-1-oxopropyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

4-[7-Chloro-2-[[[(3*R*)-1-(2,2-dimethyl-1-oxopropyl)hexahydro-2-oxo-1*H*-azepin-3-yl]amino]carbonyl]amino]-4-quinolinyl]-*N*-[(3*S*)-1-(2,2-dimethyl-1-oxopropyl)hexahydro-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-1-(methylsulfonyl)-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

(3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-carboxylic acid, methyl ester;

(3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-carboxylic acid-ethyl ester;

5 (3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-carboxylic acid,2,2,2-trifluoroethyl ester;

(3S)-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-carboxylic acid, 2-propenyl ester;

(3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-carboxylic acid phenyl ester;

10 (3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-carboxylic acid-4-fluorophenyl ester;

(3S)-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-carboxylic acid, phenylmethyl ester;

15 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-1-cyanohexahydro-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

(3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-*N*-cyclohexylhexahydro-2-oxo-1*H*-azepine-1-carboxamide;

(3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-

20 oxo-1*H*-azepine-1-carboxamide;

(2S)-2-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-6-[[[(trichloroacetyl)amino]carbonyl]amino]-hexanoic acid, methyl ester;

(3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-*N*-(2,2,2-trifluoroethyl)-1*H*-azepine-1-carboxamide;

25 (3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-*N*-cyclopropylhexahydro-2-oxo-1*H*-azepine-1-carboxamide;

(3S)-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-*N*-propyl-1*H*-azepine-1-carboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(3S)-hexahydro-2-oxo-1-(1-pyrrolidinylcarbonyl)-

30 1*H*-azepin-3-yl]-1-piperazinecarboxamide;

(3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-*N*-cyclopentylhexahydro-2-oxo-1*H*-azepine-1-carboxamide;

(3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-*N*-cyclobutylhexahydro-2-oxo-1*H*-azepine-1-carboxamide;

(3*R*)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-*N*-(4-fluorophenyl)hexahydro-2-oxo-1*H*-azepine-1-carboxamide;

5 (3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-*N*-ethylhexahydro-2-oxo-1*H*-azepine-1-carboxamide;

(3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-*N*-phenyl-1*H*-azepine-1-carboxamide;

(3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-*N*-10 (1-methylethyl)-2-oxo-1*H*-azepine-1-carboxamide;

(3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-*N*-butylhexahydro-2-oxo-1*H*-azepine-1-carboxamide;

(3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-*N*-(2-methylpropyl)-2-oxo-1*H*-azepine-1-carboxamide;

15 (3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-*N*-(1,1-dimethylethyl)hexahydro-2-oxo-1*H*-azepine-1-carboxamide;

(3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-*N,N*-dimethyl-2-oxo-1*H*-azepine-1-carboxamide;

(3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-*N*-[4-(trifluoromethyl)phenyl]-1*H*-azepine-1-carboxamide;

20 (3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-*N*-(3-fluorophenyl)hexahydro-2-oxo-1*H*-azepine-1-carboxamide;

(3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-*N*-(4-methylphenyl)-2-oxo-1*H*-azepine-1-carboxamide;

25 (3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-*N*-(2-fluorophenyl)hexahydro-2-oxo-1*H*-azepine-1-carboxamide;

(3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-*N*-(2,6-diethylphenyl)hexahydro-2-oxo-1*H*-azepine-1-carboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(3S)-hexahydro-1-[(methylamino)carbonothioyl]-

30 2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(3S)-1-[(ethylamino)carbonothioyl]hexahydro-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-1-[(1-methylethyl)amino]carbonothioyl]-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-2-oxo-1-[(2-propenylamino)carbonothioyl]-1*H*-azepin-3-yl]-1-piperazinecarboxamide;

5 (3*S*)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-*N*-methyl-2-oxo-1*H*-azepine-1-carboxamide;

(3*S*)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-*N*-(2-propenyl)-1*H*-azepine-1-carboxamide;

(3*S*)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-*N*-(2-chloroethyl)hexahydro-2-oxo-1*H*-azepine-1-carboxamide;

10 (3*R*)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-*N*-methyl-2-oxo-1*H*-azepine-1-carboxamide;

4-[3-(tert-Butoxycarbonylamino)pyrrolidin-1-yl]-7-chloroquinoline;

4-[3-(4-Fluorophenylaminocarbonylamino)pyrrolidin-1-yl]-7-chloroquinoline;

15 4-[3-[tert-Butoxycarbonyl(methyl)amino]pyrrolidin-1-yl]-7-chloroquinoline;

4-[3-[4-Fluorophenylaminocarbonyl(methyl)amino]pyrrolidin-1-yl]-7-chloroquinoline;

or enantiomers, diastereomers, salts and solvates thereof.

20 Other preferred embodiments of the present invention include:

a) A method of treating inflammation in a mammal in need thereof comprising administering to said mammal an effective anti-inflammatory amount of such compounds.

25 b) A pharmaceutical composition comprising a compound of the present invention in admixture with a pharmaceutically acceptable excipient, diluent, or carrier.

c) A method of inhibiting the binding of MIP-1 α to its receptor comprising administering a therapeutically effective amount of such compound to a mammal in need thereof.

30 d) A method of inhibiting the binding of RANTES to its receptor comprising administering a therapeutically effective amount of such compounds to a mammal in

need thereof.

e) A method of assaying compounds which modulate the activity of CCR-5 receptors comprising screening against compounds of formulae (1a) and/or (1b).

5

f) A method of inhibiting the binding of MIP-1 β to its receptor comprising administering a therapeutically effective amount of such compound to a mammal in need thereof.

g) A method of inhibiting the binding of MCP-2 to its receptor comprising
10 administering a therapeutically effective amount of such compound to a mammal in
need thereof.

The selective inhibition of CCR5 receptor mechanisms by treatment with the receptor
15 antagonists of the invention represents a novel therapeutic and preventative approach
to the treatment of a broad spectrum of inflammatory and autoimmune diseases or
conditions, in particular for the treatment of inflammatory diseases or conditions,
atherosclerosis, restenosis, and autoimmune disorders such as arthritis, Graves
disease and transplant rejection.

20

In a preferred embodiment, the disease or condition is one which is associated with
lymphocyte and/or monocyte infiltration of tissues (including recruitment and/or
accumulation in tissues), such as arthritis (e.g., rheumatoid arthritis, psoriatic arthritis,
etc.), inflammatory bowel diseases (e.g., Crohn's disease, ulcerative colitis), multiple
25 sclerosis, idiopathic pulmonary fibrosis, and graft rejection (e.g., in transplantation),
including allograft rejection or graft-versus-host disease. In addition, diseases
characterized by basophil activation and/or eosinophil recruitment, including allergic
hypersensitivity disorders such as psoriasis, asthma and allergic rhinitis can be treated
according to the present invention.

30

Other diseases and conditions that may be treated with the compounds of Formula 1
are: optic neuritis, uveitis, stroke, dermatitis, demyelinating disorders (e.g., Guillain-
Barre etc.), Grave's disease, cancer (including myeloma, breast, cervical, lymphoma,

Hodgkin's disease and Hairy Cell leukemia), endometriosis, stroke, chronic contact dermatitis, sarcoidosis, dermatomyositis, skin pemphigoid and related diseases (e.g., pemphigus vulgaris, p. foliacious, p. erythematosus), glomerulonephritides, vasculitides (e.g., necrotizing, cutaneous, and hypersensitivity vasculitis), hepatitis, 5 diabetes, systemic lupus erythematosus and myasthenia gravis.

In addition to psoriasis, other inflammatory dermatoses such as dermatitis, eczema, atopic dermatitis, allergic contact dermatitis, urticaria and reperfusion injury can also be treated.

10

It will be understood that the substituent(s) R¹, R², R³, NR⁹R⁷ and R⁴ may be at any open position on the rings of formulae (Ia) and/or (1b) to which the substituent is attached. In addition, it will be understood that there may be more than one substituent R⁴ in any given compound of formula (I), and that if there is more than one substituent 15 R⁴, that substituent may be the same or different.

The term "alkyl" is used herein at all occurrences (or a group per se or a part of a group) to mean a straight or branched chain radical of 1 to 6 carbon atoms, unless the chain length is otherwise limited, including, but not limited to methyl, ethyl, n-propyl, 20 isopropyl, n-butyl, sec-butyl, isobutyl, tert-butyl, and the like. Substituted alkyl groups may be substituted one or more times by halogen, aryl, hydroxy, R⁹-O-, amino, substituted amino, nitro, carboxy, alkoxycarbonyl, or cyano.

25 The term "cycloalkyl" is used herein at all occurrences to mean cyclic radicals, preferably of 3 to 8 carbons, including but not limited to cyclopropyl, cyclopentyl, cyclohexyl, cycloheptyl and the like. These groups can also contain one to three (as appropriate) double bonds to form the "cycloalkenyl" groups of the invention. Suitable substituents are halogen, C₁₋₆ alkyl, aryl, arylalkyl, alkylcarbonyl, hydroxy, R⁹-O-, 30 amino, substituted amino, oxo, nitro, carboxy, or cyano.

Suitable substituents on the amino groups herein can be the same or different and

include R⁹, alkyl (optionally substituted), cycloalkyl (optionally substituted), C₁₋₆ acyl, (C₁₋₆ alkoxy)carbonyl, and (C₁₋₆ alkylamino)carbonyl . Typical substituents include OH, C₁₋₆ alkoxy, C₁₋₆ alkyl, and R⁹ aminocarbonyloxy.

- 5 The terms "halo" or "halogen" are used interchangeably herein at all occurrences to mean radicals derived from the elements chlorine, fluorine, iodine and bromine. "Halogenated " is analogous and refers to a degree of halogen substitutions from single to full (per) substitution.
- 10 The terms "aryl" or "Ar--" are used herein at all occurrences to mean 5-10 membered (fused or connected) aromatic ring(s) or ring systems which may include bi- or tri-cyclic systems. Representative examples include, but are not limited to, phenyl and naphthyl. Substituted aryl groups may be substituted one or more times by halogen, C₁₋₆ alkyl, hydroxy, alkoxy, e.g, methoxy, amino, substituted amino, nitro, trifluoromethyl,
- 15 carboxy, acyl, or cyano.

The term "heterocyclyl" (heterocyclic or heterocycle or heterocyclo) refers to a stable 3- to 15-membered ring radical which consists of carbon atoms and from one to five heteroatoms selected from the group consisting of nitrogen, oxygen and sulfur. For purposes of this invention, the heterocyclyl radical may be a monocyclic, bicyclic or tricyclic ring system, which may include fused or bridged ring systems; and the nitrogen, carbon or sulfur atoms in the heterocyclyl radical may be optionally oxidized; the nitrogen atom may be optionally quaternized; and the heterocyclyl radical may be partially or fully saturated or aromatic (i.e., heteroaryl). The heterocyclyl radical may be attached to the main structure at any heteroatom or carbon atom which results in the creation of a stable compound. Examples of such heterocyclyl radicals include, but are not limited to Examples of heterocyclic groups include, e.g., piperidinyl, piperazinyl, oxopiperazinyl, oxopiperidinyl, oxopyrrolodinyl, oxoazepinyl, azepinyl, pyrrolyl, piperidonyl, pyrrolidinyl, pyrazolyl, pyrazolidinyl, imidazolyl, imidazolinyl, imidazolidinyl, pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, oxazolyl, oxazolidinyl, isoxazolyl, isoxazolidinyl, morpholinyl, thiazolyl, thiophenyl, oxazolyl, thiazolidinyl, isothiazolyl, quinuclidinyl, isothiazolidinyl, indolyl, quinolinyl, isoquinolinyl, benzimidazolyl, thiadiazoyl, benzopyranyl, benzothiazolyl, benzoxazolyl, furyl,

tetrahydrofuryl, tetrahydropyranyl, tetrazole, thienyl, benzothienyl, thiamorpholinyl, thiamorpholinyl sulfoxide, thiamorpholinyl sulfone, benzimidazolyl, benzofuranyl, benzothiophenyl, benzoxazolyl, furyl, imidazolyl, indolyl, isoquinolyl, isothiazolyl, isoxazolyl, morpholinyl, oxadiazolyl, oxazolyl, piperidyl, pyrazolyl, pyrimidyl, pyrrolyl,
 5 quinolyl, tetrahydroisoquinolyl, tetrahydroquinolyl, tetrahydrothienyl, tetrazolyl, thiadiazolyl, thiazolyl, triazolyl; pyrrolidine, piperidine, pyridine, tetrahydrofuran, thiophene, oxazole, thiazole, imidazole, benzothiazole, benzoxazole, benzimidazole, selenazole, benzoselenazole, tellurazole, triazole, benzotriazole, tetrazole, benzofuranyl, benzothienyl, oxadiazole, or thiadiazole rings. A preferred heterocyclic
 10 group is hexahydroazepinyl.

Heteroaryl refers to a fully unsaturated heterocycle having one or two rings and a total number of 5 to 10 ring atoms wherein at least one of the ring atoms is a heteroatom. Preferably, the heteroaryl group contains 1 to 3 hetero atoms which are selected from
 15 N, O and S. Suitable heteroaryl groups include but are not limited to furyl, thienyl, pyrrolyl, pyrazolyl, imidazolyl, triazolyl, dithialyl, oxathialyl, isoxazolyl, oxazolyl, thiazolyl, isothiazolyl, oxadiazolyl, oxatriazolyl, dioxazolyl, oxathiazolyl, thiadiazolyl, pyridyl, pyridazinyl, pyrimidinyl, pyrazinyl, triazinyl, oxazinyl, isoxazinyl, oxathiazinyl, oxadiazinyl, benzofuranyl, isobenzofuranyl, thionaphthenyl, isothionaphthenyl, indolyl,
 20 isoindolyl, indazolyl, benzisoxazolyl, benzoxazolyl, benzthiazolyl, benzisothiazolyl, purinyl, benzopyranyl, quinolinyl, isoquinolinyl, cinnolinyl, quinazolinyl, naphthyridinyl, and benzoxazinyl, e.g., 2-thienyl, 3-thienyl, 2-, 3- or 4-pyridyl, 2-, 3-, 4-, 5-, 6-, 7- or 8-quinolinyl, and 1-, 3-, 4-, 5-, 6-, 7- or 8-isoquinolinyl.
 25 Substituted heterocyclyl refers to the heterocyclyl groups described above which are substituted in one or more places by R⁹, COR⁹, -SO₂-R⁷, halogen, aryl, alkyl, alkoxy, methylene, cyano, trifluoromethyl, carboxy, nitro, oxo, amino, alkoxycarbonyl, aminocarbonyl, substituted aminocarbonyl, or substituted amino.
 30 Alkoxy groups means alkyl-O- groups in which the alkyl portion (substituted or unsubstituted) is in accordance with the previous discussion. Suitable alkoxy groups are methoxy, ethoxy, propoxy and butoxy.

The term "oxo" refers to the group =O, and the term "thioxo" refers to the group =S.

Acyl refers to alkanoyl radicals having 1 to 6 carbon atoms in which the alkyl portion can be substituted as defined above.

5

It will be understood throughout that the optional substituents are selected independently from one another.

10 The term "CCR5 mediated disease state" is used herein at all occurrences to mean any disease state which is affected or modulated by CCR5.

15 Some of the compounds of Formula 1a and 1b and related compounds are capable of forming both pharmaceutically acceptable acid addition and/or base salts. All of these forms are within the scope of the present invention, as are diastereomers and enantiomers.

The optical isomers can be obtained by resolution of the racemic mixtures according to conventional processes, for example, by the formation of diastereoisomeric salts using an optically active acid or base, or by formation of covalent diastereomers. Examples 20 of appropriate acids are tartaric, diacetyl tartaric, dibenzoyl tartaric, ditoluoyl tartaric and camphorsulfonic acid. Mixtures of diastereoisomers can be separated into their individual diastereomers on the basis of their physical and/or chemical differences by methods known to those skilled in the art, for example, by chromatography or fractional crystallization. The optically active bases or acids are then liberated from 25 the separated diastereomeric salts. A different process for separation of optical isomers involves use of chiral chromatography (e.g., chiral HPLC columns), with or without conventional derivitization, optimally chosen to maximize the separation of the enantiomers. Suitable chiral HPLC columns are manufactured by Diacel, e.g., Chiracel OD and Chiracel OJ, among many others, all routinely selectable. Enzymatic 30 separations, with or without derivitization, are also useful. The optically active compounds of formula I can likewise be obtained by utilizing optically active starting materials.

Compounds of the present invention can exist as salts (preferably pharmaceutically acceptable salts). Pharmaceutically acceptable acid addition salts of the compounds of Formulae (1a) and (1b) include salts derived from nontoxic inorganic acids such as hydrochloric, nitric, phosphoric, sulfuric, hydrobromic, hydriodic, hydrofluoric, 5 phosphorous, and the like, as well as the salts derived from nontoxic organic acids, such as aliphatic mono- and dicarboxylic acids, 2-phenyl-substituted alkanoic acids, hydroxy alkanoic acids, alkanedioic acids, aromatic acids, aliphatic and aromatic sulfonic acids, etc. Such salts thus include sulfate, pyrosulfate, bisulfate, sulfite, bisulfite, nitrate, phosphate, monohydrogenphosphate, dihydrogenphosphate, 10 metaphosphate, pyrophosphate, chloride, bromide, iodide, acetate, trifluoroacetate, propionate, caprylate, isobutyrate, oxalate, malonate, succinate, suberate, sebacate, fumarate, maleate, mandelate, benzoate, chlorobenzoate, methylbenzoate, dinitrobenzoate, phthalate, benzenesulfonate, toluenesulfonate, phenylacetate, citrate, lactate, maleate, tartrate, methanesulfonate, and the like. Also contemplated are salts 15 of amino acids such as arginate and the like and gluconate, galacturonate (see, for example, Berge S. M. et al., "Pharmaceutical Salts," J. Pharma. Sci., 1977;66:1). The acid addition salts of basic compounds can be prepared by contacting the free base form with a sufficient amount of the desired acid to produce the salt in the conventional manner. The free base form may be regenerated by contacting the salt 20 form with a base and isolating the free base in the conventional manner. The free base forms can differ from their respective salt forms somewhat in certain physical properties such as solubility in polar solvents.

Pharmaceutically acceptable base addition salts can be formed with metals or amines, 25 such as alkali and alkaline earth metals or organic amines. Examples of such metals used as cations are sodium, potassium, magnesium, calcium, and the like. Examples of suitable amines are N,N'-dibenzylethylenediamine, chloroprocaine, choline, diethanolamine, dicyclohexylamine, ethylenediamine, N-methylglucamine, and procaine (see Berge, Supra, 1977).

The base addition salts of said acidic compounds can be prepared by contacting the free acid form with a sufficient amount of the desired base to produce the salt in the conventional manner. The free acid form may be regenerated by contacting the salt 30

form with an acid and isolating the free acid in the conventional manner. The free acid forms can differ from their respective salt forms somewhat in certain physical properties such as solubility in polar solvents.

5 Certain of the compounds of the present invention can exist in unsolvated forms as well as solvated forms, including hydrated forms and are intended to be encompassed within the scope of the present invention.

10 Certain of the compounds of the present invention possess one or more chiral centers and each center may exist in the R(D) or S(L) configuration. The present invention includes all diastereomeric, enantiomeric and epimeric forms as well as all mixtures thereof.

15 The activity of compounds of the present invention can be assessed using suitable assays, such as receptor binding assays and chemotaxis assays. For example, as described in the Example Section, antagonist compounds of the present invention have been identified utilizing a CCR5 Receptor MIP1 α SPA binding assay and have been found to exhibit IC₅₀ values ranging from 0.005 μ M to 38 μ M. Such values are indicative of the intrinsic activity of the compounds in use as modulators of chemokine 20 receptor activity. There are numerous other such screening assays known to those skilled in the art which may be used to determine the CCR5 receptor antagonistic activity of the compounds of the present invention. One such screening technique is described in PCT WO 92/01810. Another assay, for example, may be employed for screening a receptor antagonist by contacting melanophore cells which encode the 25 CC-CKR5 receptor with both a chemokine agonist and a compound to be screened. Inhibition of the signal generated by the ligand indicates that a compound is a potential antagonist for the receptor, i.e., inhibits activation of the receptor.

30 Other screening techniques include the use of cells which express the CC-CKR5 receptor (for example, transfected CHO cells, RBL-2 cells or other mammalian cells) in a system which measures extracellular pH changes caused by receptor activation, for example, as described in Science, volume 246, pages 181-296 (October 1989), herein incorporated by reference. For example, potential antagonists may be contacted with a

cell which expresses the CC-CKR5 receptor and a second messenger response, e.g. signal transduction or pH changes, or making use of a reporter gene system, for example luciferase, may be measured to determine whether the potential antagonist is effective.

5

Another such screening technique involves introducing mRNA encoding the CC-CKR5 receptor into mammalian cells (Xenopus oocytes, RBL-2 or other mammalian cells) to transiently express the receptor. The cells with the expressed receptor may then be contacted in the case of antagonist screening with a chemokine agonist and a

10 compound to be screened, followed by detection of inhibition of a calcium or cAMP signal.

Another screening technique involves expressing the CC-CKR5 receptor in which the receptor is linked to a phospholipase C or D. As representative examples of such cells, 15 there may be mentioned endothelial cells, smooth muscle cells, embryonic kidney cells, etc. The screening for an antagonist may be accomplished as hereinabove described by detecting inhibition of activation of the receptor from the phospholipase second signal.

20 Another method involves screening for CC-CKR5 receptor inhibitors by determining inhibition of binding of labeled chemokine ligands to cells or membranes which have the receptor on the surface thereof. Such a method involves transfecting a eukaryotic cell, such as CHO or RBL-2 cell, with DNA encoding the CC-CKR5 receptor such that the cell expresses the receptor on its surface and contacting the cell with a potential 25 antagonist in the presence of a labeled form of chemokine ligand. The chemokine ligand can be labeled, e.g., by radioactivity. The amount of labeled ligand bound to the receptors is measured, e.g., by measuring radioactivity associated with transfected cells or membrane from these cells. If the potential antagonist binds to the receptor, as determined by a reduction of labeled ligand which binds to the receptors, the binding 30 of labeled ligand to the receptor is inhibited.

Another method involves screening for CC-CKR5 inhibitors by determining inhibition or stimulation of CC-CKR5-mediated cAMP and/or adenylate cyclase accumulation or

dimunition. Such a method involves transfecting a eukaryotic cell, such as CHO or RBL-2 cell, with CC-CKR5 receptor to express the receptor on the cell surface. The cell is then exposed to potential antagonists in the presence of chemokine agonist.

The amount of cAMP accumulation is then measured. If the potential antagonist binds
5 the receptor, and thus inhibits CC-CKR5 binding, the levels of CC-CKR5-mediated cAMP, or adenylate cyclase, activity will be reduced or increased.

Another such screening technique is described in USP 5,928,881, which provides a method for determining whether a ligand not known to be capable of binding to the CC-CKR5 receptor can bind to such receptor which comprises contacting a
10 mammalian cell which expresses the CC-CKR5 receptor with a chemokine ligand such as RANTES under conditions permitting binding of ligands to the CC-CKR5 receptor, detecting the presence of a ligand which binds to the receptor and thereby determining whether the ligand binds to the CC-CKR5 receptor.

15 Another method involves screening of CCR5 expressed in yeast cells that have been modified to couple receptor activation by chemokine ligands to the yeast phermone response.

Another method involves screening of a CCR5 receptor that has been genetically
20 modified so that it exhibits constitutive activity in either mammalian or yeast cells. In this case an activating chemokine ligand is not required and the nature of the receptor modulation (agonist, inverse agonism, antagonism, etc.) can be specifically addressed.

25 The antagonists of the present invention bind to the CC-CKR5 receptor, making it inaccessible to ligands such that normal biological activity is prevented. They may be administered to a mammal in need of treatment of CCR5 mediated disease states. Thus, the active ingredient may be administered in the mammal using conventional course of treatment determination tests.

30 The subject treated in the methods above is a mammal, preferably a human being, male or female, in whom modulation of chemokine receptor activity is desired. "Modulation" as used herein is intended to encompass antagonism, agonism, partial

antagonism, inverse agonism and/or partial agonism. In a preferred aspect of the present invention, modulation refers to antagonism or inverse agonism of chemokine receptor activity, since the compounds of the invention are antagonists or inverse agonists.

5

Combined therapy to modulate chemokine receptor activity and thereby prevent and treat inflammatory and immunoregulatory disorders and diseases, including endometriosis, asthma and allergic diseases, as well as autoimmune pathologies such as rheumatoid arthritis, Graves' disease and atherosclerosis, and those pathologies noted above is illustrated by the combination of the compounds of this invention and other compounds which are known for such utilities.

For example, in the treatment or prevention of inflammation, the present compounds may be used in conjunction with an antiinflammatory or analgesic agent such as an opiate agonist, a lipoxygenase inhibitor, such as an inhibitor of 5-lipoxygenase, a cyclooxygenase inhibitor, such as a cyclooxygenase-2 inhibitor, an interleukin inhibitor, such as an interleukin-1 inhibitor, an NMDA antagonist, an inhibitor of nitric oxide or an inhibitor of the synthesis of nitric oxide, a non-steroidal antiinflammatory agent, or a cytokine-suppressing antiinflammatory agent, for example with a compound such as acetaminophen, aspirin, codeine, fentanyl, ibuprofen, indomethacin, ketorolac, morphine, naproxen, phenacetin, piroxicam, a steroidal analgesic, sufentanil, sunlindac, tenidap, and the like. Similarly, the instant compounds may be administered with a pain reliever; a potentiator such as caffeine, an H₂-antagonist, simethicone, aluminum or magnesium hydroxide; a decongestant such as phenylephrine, phenylpropanolamine, pseudophedrine, oxymetazoline, epinephrine, naphazoline, xylometazoline, propylhexedrine, or levo-desoxy-ephedrine; an antiitussive such as codeine, hydrocodone, caramiphen, carbetapentane, or dextromethorphan; a diuretic; and a sedating or non-sedating antihistamine. Likewise, compounds of the present invention may be used in combination with other drugs that are used in the treatment/prevention/suppression or amelioration of the diseases or conditions for which compounds of the present invention are useful. Such other drugs may be administered, by a route and in an amount commonly used therefore, contemporaneously or sequentially with a compound of the present invention. When a

compound of the present invention is used contemporaneously with one or more other drugs, a pharmaceutical composition containing such other drugs in addition to the compound of the present invention is preferred.

5 Accordingly, the pharmaceutical compositions of the present invention include those that also contain one or more other active ingredients, in addition to a compound of the present invention. Examples of other active ingredients that may be combined with a compound of the present invention, either administered separately or in the same pharmaceutical compositions, include, but are not limited to: (a) VLA-4 antagonists
10 such as those described in U.S. Pat. No. 5,510,332, (b) steroids such as beclomethasone, methylprednisolone, betamethasone, prednisone, dexamethasone, and hydrocortisone; (c) immunosuppressants such as cyclosporin, tacrolimus, rapamycin and other FK-506 type immunosuppressants; (d) antihistamines (H1-histamine antagonists) such as bromopheniramine, chlorpheniramine,
15 dexchlorpheniramine, triprolidine, clemastine, diphenhydramine, diphenylpyraline, tripelennamine, hydroxyzine, methdilazine, promethazine, trimeprazine, azatadine, cyproheptadine, antazoline, pheniramine pyrilamine, astemizole, terfenadine, loratadine, cetirizine, fexofenadine, descarboethoxyloratadine, and the like; (e) non-steroidal anti-asthmatics such as beta 2-agonists (terbutaline, metaproterenol,
20 fenoterol, isoetharine, albuterol, bitolterol, and pirbuterol), theophylline, cromolyn sodium, atropine, ipratropium bromide, leukotriene antagonists (zafirlukast, montelukast, pranlukast, iralukast, pobilukast, SKB-106,203), leukotriene biosynthesis inhibitors (zileuton, BAY-1005); (f) non-steroidal antiinflammatory agents (NSAIDs)
25 such as propionic acid derivatives (alminoprofen, benoxaprofen, bucloxic acid, caiprofen, fenbufen, fenoprofen, fluprofen, flurbiprofen, ibuprofen, indoprofen, ketoprofen, mioprofen, naproxen, oxaprozin, pirprofen, pranoprofen, suprofen, tiaprofenic acid, and tioxaprofen), acetic acid derivatives (indomethacin, acemetacin, alclofenac, clidanac, diclofenac, fenclofenac, fenclozic acid, fentiazac, furofenac, ibufenac, isoxepac, oxpainac, sulindac, tiopinac, tolmetin, zidometacin, and zomepirac),
30 fenamic acid derivatives (flufenamic acid, meclofenamic acid, mefenamic acid, niflumic acid and tolfenamic acid), biphenylcarboxylic acid derivatives (diflunisal and flufenisal), oxicams (isoxicam, piroxicam, sudoxicam and tenoxicam), salicylates (acetyl salicylic acid, sulfasalazine) and the pyrazolones (apazone, bezpiperylon, feprazone,

mofebutazone, oxyphenbutazone, phenylbutazone); (g) cyclooxygenase-2 (COX-2) inhibitors; (h) inhibitors of phosphodiesterase type IV (PDE-IV); (i) other antagonists of the chemokine receptors, CCR1, CCR2, CCR3, CCR4, CCR8, CCR10, CXCR1, CXCR2, CXCR3, and CX3CR1; (j) cholesterol lowering agents such as HMG-CoA reductase inhibitors (lovastatin, simvastatin and pravastatin, fluvastatin, atorvastatin, and other statins), sequestrants (cholestyramine and colestipol), nicotinic acid, fenofibric acid derivatives (gemfibrozil, clofibrate, fenofibrate and benzafibrate), and probucol; (k) anti-diabetic agents such as insulin, sulfonylureas, biguanides (metformin), α -glucosidase inhibitors (acarbose) and glitazones (troglitazone and pioglitazone); (l) preparations for multiple sclerosis such as interferon beta 1b (Betaseron), interferon-beta 1a (Avonex), glatiramer acetate (Copaxone), azathioprine (Imurek, Imuran), mitoxantrone (Novatrone), IgG, cyclophosphamide; (m) agents for dermatological conditions such as a lubricant, a keritolytic agent, vitamin D3 derivative, PUVA, anthralin; (n) other compounds such as 5-aminosalicylic acid and prodrugs thereof, antimetabolites such as azathioprine and 6-mercaptopurine, and cytotoxic cancer chemotherapeutic agents; (o) estrogen receptor agonists such as estriol, estradiol, or any synthetic ER α or ER β agonist; (p) IL-2 receptor antagonists such as an anti-IL-2 receptor antibody (Zenapax); (q) IL-12 receptor antagonists; (r) T-helper cytokines such as IL-4 or IL-10; (s) anti-CD52 antibody (Campath).

20

The weight ratio of the compound of the present invention to the second active ingredient may be varied and will depend upon the effective dose of each ingredient. Generally, an effective dose of each will be used. Thus, for example, when a compound of the present invention is combined with an NSAID the weight ratio of the compound of the present invention to the NSAID will generally range from about 1000:1 to about 1:1000, preferably about 200:1 to about 1:200. Combinations of a compound of the present invention and other active ingredients will generally also be within the aforementioned range, but in each case, an effective dose of each active ingredient should be used.

25
30

The compounds of the present invention may be administered by oral, parenteral (e.g., intramuscular, intraperitoneal, intravenous, ICV, intracisternal injection or infusion, subcutaneous injection, or implant), by inhalation spray, nasal, vaginal, rectal,

sublingual, or topical routes of administration and may be formulated, alone or together, in suitable dosage unit formulations containing conventional non-toxic pharmaceutically acceptable carriers, adjuvants and vehicles appropriate for each route of administration. The compounds of the invention are effective for use in
5 primates, such as humans, as well as for the treatment of warm-blooded animals such as mice, rats, horses, cattle, sheep, dogs, cats, monkeys, guinea pigs, other bovine, ovine, equine, canine, feline, rodent or murine species. However, the compounds of the invention are also effective for use in other species, such as avian species (e.g., chickens).

10

The pharmaceutical compositions for the administration of the compounds of this invention may conveniently be presented in dosage unit form and may be prepared by any of the methods well known in the art of pharmacy. All methods include the step of bringing the active ingredient into association with the carrier which constitutes one or
15 more accessory ingredients. In general, the pharmaceutical compositions are prepared by uniformly and intimately bringing the active ingredient into association with a liquid carrier or a finely divided solid carrier or both, and then, if necessary, shaping the product into the desired formulation. In the pharmaceutical composition the active object compound is included in an amount sufficient to produce the desired effect upon
20 the process or condition of diseases.

The pharmaceutical compositions containing the active ingredient may be in a form suitable for oral use, for example, as tablets, troches, lozenges, aqueous or oily suspensions, dispersible powders or granules, emulsions, hard or soft capsules, or
25 syrups or elixirs. Compositions intended for oral use may be prepared according to any method known to the art for the manufacture of pharmaceutical compositions and such compositions may contain one or more agents selected from the group consisting of sweetening agents, flavoring agents, coloring agents and preserving agents in order to provide pharmaceutically elegant and palatable preparations. Tablets contain the
30 active ingredient in admixture with non-toxic pharmaceutically acceptable excipients which are suitable for the manufacture of tablets. These excipients may be for example, inert diluents, such as calcium carbonate, sodium carbonate, lactose, calcium phosphate or sodium phosphate; granulating and disintegrating agents, for

example, corn starch, or alginic acid; binding agents, for example starch, gelatin or acacia, and lubricating agents, for example magnesium stearate, stearic acid or talc. The tablets may be uncoated or they may be coated by known techniques to delay disintegration and absorption in the gastrointestinal tract and thereby provide a

5 sustained action over a longer period. For example, a time delay material such as glyceryl monostearate or glyceryl distearate may be employed. They may also be coated by the techniques described in the U.S. Pat. Nos. 4,256,108; 4,166,452; and 4,265,874 to form osmotic therapeutic tablets for control release.

10 Formulations for oral use may also be presented as hard gelatin capsules wherein the active ingredient is mixed with an inert solid diluent, for example, calcium carbonate, calcium phosphate or kaolin, or as soft gelatin capsules wherein the active ingredient is mixed with water or an oil medium, for example peanut oil, liquid paraffin, or olive oil.

15 Aqueous suspensions contain the active materials in admixture with excipients suitable for the manufacture of aqueous suspensions. Such excipients are suspending agents, for example sodium carboxymethylcellulose, methylcellulose, hydroxy-propylmethylcellulose, sodium alginate, polyvinyl-pyrrolidone, gum tragacanth and gum acacia; dispersing or wetting agents may be a naturally-occurring phosphatide,

20 for example lecithin, or condensation products of an alkylene oxide with fatty acids, for example polyoxyethylene stearate, or condensation products of ethylene oxide with long chain aliphatic alcohols, for example heptadecaethyleneoxycetanol, or condensation products of ethylene oxide with partial esters derived from fatty acids and a hexitol such as polyoxyethylene sorbitol monooleate, or condensation products

25 of ethylene oxide with partial esters derived from fatty acids and hexitol anhydrides, for example polyethylene sorbitan monooleate. The aqueous suspensions may also contain one or more preservatives, for example ethyl, or n-propyl, p-hydroxybenzoate, one or more coloring agents, one or more flavoring agents, and one or more sweetening agents, such as sucrose or saccharin.

30 Oily suspensions may be formulated by suspending the active ingredient in a vegetable oil, for example arachis oil, olive oil, sesame oil or coconut oil, or in a mineral oil such as liquid paraffin. The oily suspensions may contain a thickening

agent, for example beeswax, hard paraffin or cetyl alcohol. Sweetening agents such as those set forth above, and flavoring agents may be added to provide a palatable oral preparation. These compositions may be preserved by the addition of an antioxidant such as ascorbic acid.

5

Dispersible powders and granules suitable for preparation of an aqueous suspension by the addition of water provide the active ingredient in admixture with a dispersing or wetting agent, suspending agent and one or more preservatives. Suitable dispersing or wetting agents and suspending agents are exemplified by those already mentioned 10 above. Additional excipients, for example sweetening, flavoring and coloring agents, may also be present.

The pharmaceutical compositions of the invention may also be in the form of oil-in-water emulsions. The oily phase may be a vegetable oil, for example olive oil or 15 arachis oil, or a mineral oil, for example liquid paraffin or mixtures of these. Suitable emulsifying agents may be naturally-occurring gums, for example gum acacia or gum tragacanth, naturally-occurring phosphatides, for example soy bean, lecithin, and esters or partial esters derived from fatty acids and hexitol anhydrides, for example sorbitan monooleate, and condensation products of the said partial esters with 20 ethylene oxide, for example polyoxyethylene sorbitan monooleate. The emulsions may also contain sweetening and flavoring agents.

Syrups and elixirs may be formulated with sweetening agents, for example glycerol, propylene glycol, sorbitol or sucrose. Such formulations may also contain a demulcent, 25 a preservative and flavoring and coloring agents.

The pharmaceutical compositions may be in the form of a sterile injectable aqueous or oleagenous suspension. This suspension may be formulated according to the known art using those suitable dispersing or wetting agents and suspending agents which 30 have been mentioned above. The sterile injectable preparation may also be a sterile injectable solution or suspension in a non-toxic parenterally-acceptable diluent or solvent, for example as a solution in 1,3-butane diol. Among the acceptable vehicles and solvents that may be employed are water, Ringer's solution and isotonic sodium

chloride solution. In addition, sterile, fixed oils are conventionally employed as a solvent or suspending medium. For this purpose any bland fixed oil may be employed including synthetic mono- or diglycerides. In addition, fatty acids such as oleic acid find use in the preparation of injectables.

5

The compounds of the present invention may also be administered in the form of suppositories for rectal administration of the drug. These compositions can be prepared by mixing the drug with a suitable non-irritating excipient which is solid at ordinary temperatures but liquid at the rectal temperature and will therefore melt in the rectum to release the drug. Such materials are cocoa butter and polyethylene glycols.

For topical use, creams, ointments, jellies, solutions or suspensions, etc., containing the compounds of the present invention are employed. (For purposes of this application, topical application shall include mouthwashes and gargles.) The pharmaceutical composition and method of the present invention may further comprise other therapeutically active compounds as noted herein which are usually applied in the treatment of the above mentioned pathological conditions.

The compounds of the invention, or their pharmaceutically acceptable salts, are administered in a therapeutically effective amount which will vary depending upon a variety of factors including the activity of the specific compound employed; the metabolic stability and length of action of the compound; the age, body weight, general health, sex, and diet of the patient; the mode and time of administration; the rate of excretion; the drug combination; the severity of the particular disease-states; and the host undergoing therapy. Generally, a therapeutically effective daily dose is from about 0.14 mg to about 14.3 mg/kg of body weight per day of a compound of the invention, or a pharmaceutically acceptable salt thereof; preferably, from about 0.7 mg to about 10 mg/kg of body weight per day; and most preferably, from about 1.4 mg to about 7.2 mg/kg of body weight per day. For example, for administration to a 70 kg person, the dosage range would be from about 10 mg to about 1.0 gram per day of a compound of the invention, or a pharmaceutically acceptable salt thereof, preferably from about 50 mg to about 700 mg per day, and most preferably from about 100 mg to about 500 mg per day.

The compounds may be administered on a regimen of 1 to 4 times per day, preferably

once or twice per day.

It will be understood, however, that the specific dose level and frequency of dosage for any particular patient may be varied and will depend upon a variety of factors including

5 the activity of the specific compound employed, the metabolic stability and length of action of that compound, the age, body weight, general health, sex, diet, mode and time of administration, rate of excretion, drug combination, the severity of the particular condition, and the host undergoing therapy.

10 Having described the invention, the following examples are given to illustrate specific applications of the invention including the best mode now known to perform the invention. These specific examples are not intended to limit the scope of the invention described in this application. Without further elaboration, it is believed that one skilled in the art can, using the preceding description, utilize the present invention to its fullest extent. The following preferred specific embodiments are, therefore, to be construed 15 as merely illustrative, and not limitative of the remainder of the disclosure in any way whatsoever.

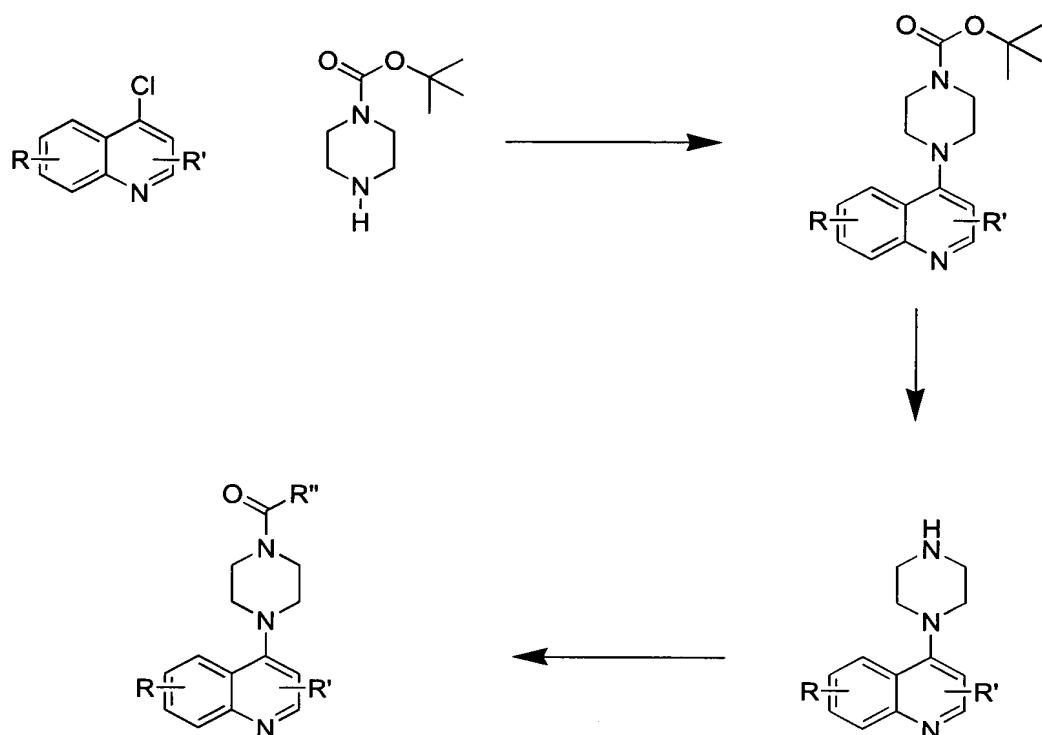
In the foregoing and in the following examples, all temperatures are set forth
20 uncorrected in degrees Celsius; and, unless otherwise indicated, all parts and percentages are by weight.

The entire disclosures of all applications, patents and publications, cited above or below, is hereby incorporated by reference.

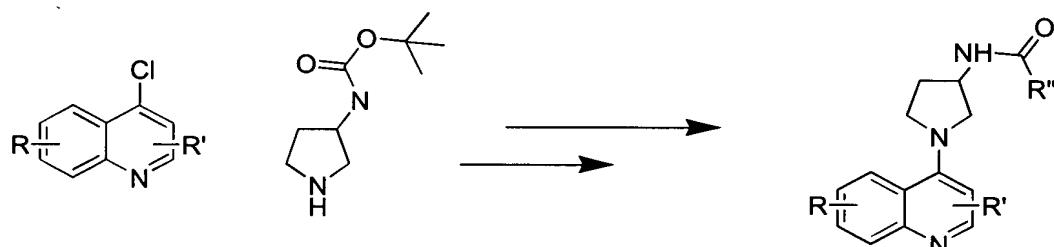
25 **General Methods of Preparation**

Specifically, the compounds of the invention are prepared according to the following general methods and schemes:

General Scheme for preparing 4-(piperazin-1-yl)quinolines



General Scheme for preparing 4-(pyrrolidin-1-yl)quinolines



Method A.

5 A 4-chloroquinoline is heated with 1-*tert*-butoxycarbonylpiperazine at 150 °C for 2 h. The reaction mixture is diluted with EtOAc, washed with water, dried (MgSO_4), and concentrated. The residue is dissolved in CH_2Cl_2 and treated with trifluoroacetic acid for 1 h. The mixture is concentrated, diluted with EtOAc, washed with sat. NaHCO_3 , dried (MgSO_4), and concentrated *in vacuo*.

10

Method B

A 4-chloroquinoline and piperazine are heated either in EtOH at reflux or in *n*-butanol at 150 °C in a sealed tube for 18 h. The reaction mixture is concentrated, and the residue dissolved in EtOAc. After washing with water, and drying (MgSO_4), the solvent
15 was removed *in vacuo*.

General Method for Reacting 4-(Piperazin-1-yl)quinolines with Isocyanates:***Method C***

Iso(thio)cyanate is added to a suspension of a 4-(piperazin-1-yl)quinoline (and
 5 optionally triethylamine) in CH₂Cl₂ or THF. The reaction mixture is concentrated after 1 h, and the residue is purified either by recrystallization or column chromatography.

Method D

Amine, nitrophenyl chloroformate, and triethylamine are stirred either in EtOAc or CH₂Cl₂ for ½ - 1 h. In case of CH₂Cl₂ solutions the solvent is removed and the residue
 10 suspended in EtOAc. A 4-(piperazin-1-yl)quinoline is added, and the reaction mixture is heated at reflux for 1 - 2 h. After filtration the filtrate is concentrated and the residue purified by column chromatography.

General Method for Reacting 2-Quinolones with Amines:***Method E***

Sodium hydride is added to a solution of a 2-quinolone in DMSO and stirred at 40 °C for 1 h, when *N*-phenyl (trifluoromethylsulfon)imide is added. After an hour amine is added, and the reaction mixture is heated in a sealed tube at 150 °C for 15 h. The mixture is diluted with EtOAc, washed with water and brine, dried (NaSO₄), and
 20 concentrated. The residue is purified by column chromatography with CH₂Cl₂-MeOH.

EXAMPLES**Example 1****Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline**

Preparation of 7-Chloro-4-(piperazin-1-yl)quinoline. 4,7-Dichloroquinoline (9.9 g, 50 mmol) and piperazine (43.1 g, 500 mmol) in EtOH (200 mL) are reacted according to
 30 method B yielding 8.10 g of the product.

¹H NMR (CDCl₃) δ 3.17 (m, 8H), 6.82 (d, 1H), 7.40 (dd, 1H), 7.93 (d, 1H), 8.01 (d, 1H), 8.69 (d, 1H).

Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline

7-Chloro-4-(piperazin-1-yl)quinoline (0.89 g, 3.6 mmol) and 4-fluorophenyl isocyanate (409 µL, 3.6 mmol) in THF (20 mL) are reacted according to method C yielding 0.71 g

5 of the product as a colorless solid after column chromatography with hexane-EtOAc.

¹H NMR ([D]₆-DMSO), δ 3.18 (m, 4H), 3.71 (m, 4H), 7.04 (d, 1H), 7.06 (t, 2H), 7.46 (dd, 2H), 7.56 (dd, 1H), 7.98 (d, 1H), 8.09 (d, 1H), 8.68 (br. s, 1H), 8.72 (d, 1H).

Example 2

10 **Preparation of 6-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline**

Preparation of 6-Chloro-4-(piperazin-1-yl)quinoline

4,6-Dichloroquinoline (De, D., Byers L.D., Krogstad, D.J. *J. Heterocyclic Chem.* **1997**, 15 34, 315) (0.34 g, 1.7 mmol) and 1-*tert*-butoxycarbonylpiperazine (1.58 g, 8.5 mmol) are reacted according to method A. The Boc group is cleaved with trifluoroacetic acid (5 mL) in CH₂Cl₂ (5 mL) giving 0.25 g of the product.

¹H NMR (CDCl₃), δ 3.13 (s, 8H), 6.91 (d, 1H), 7.53 (dd, 1H), 7.92 (d, 1H), 7.94 (d, 1H), 8.66 (d, 1H).

20

Preparation of 6-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline

6-Chloro-4-(piperazin-1-yl)quinoline (0.25 g, 1.0 mmol), 4-fluorophenyl isocyanate (170 µL, 1.5 mmol), and DMAP (2 mg) in THF (10 mL) are reacted according to method C 25 yielding the product as a colorless solid after column chromatography with hexane-EtOAc. ¹H NMR ([D]₆-DMSO), δ 3.17 (m, 4H), 3.75 (m, 4H), 7.08 (t, 2H), 7.10 (d, 1H), 7.46 (dd, 2H), 7.72 (dd, 1H), 7.98 (d, 1H), 8.01 (d, 1H), 8.68 (br. s, 1H), 8.72 (d, 1H).

Example 3

30 **Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-methylquinoline**

Preparation of 7-Chloro-2-methyl-4-(piperazin-1-yl)quinoline

4,7-Dichloro-2-methylquinoline (De, D., Byers L.D., Krogstad, D.J. *J. Heterocyclic Chem.* **1997**, *34*, 315) (453 mg, 2.1 mmol) and 1-*tert*-butoxycarbonylpiperazine (2.05 g, 11.0 mmol) are reacted according to method A. The Boc group is cleaved with

5 trifluoroacetic acid (5 mL) in CH₂Cl₂ (5 mL) giving 0.26 g of the piperazine.

¹H NMR (CDCl₃), δ 2.62 (s, 3H), 3.12 (s, 8H), 6.67 (s, 1H), 7.30 (dd, 1H), 7.83 (d, 1H), 7.90 (d, 1H).

Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-

10 *methylquinoline*

7-Chloro-2-methyl-4-(piperazin-1-yl)quinoline (0.26 g), 4-fluorophenyl isocyanate (114 μL, 1.0 mmol), and DMAP (2 mg) in THF (10 mL) are reacted according to method C yielding the product as a colorless solid after column chromatography with hexane-

15 EtOAc. ¹H NMR ([D]₆-DMSO), δ 2.58 (s, 3H), 3.17 (m, 4H), 3.71 (m, 4H), 6.94 (s, 1H), 7.06 (t, 2H), 7.46 (m, 3H), 7.87 (d, 1H), 8.01 (d, 1H), 8.68 (br. s, 1H).

Example 4*Preparation of 5-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-*

20 *methylquinoline*

Preparation of 5-Chloro-2-methyl-4-(piperazin-1-yl)quinoline

4,5-Dichloro-2-methylquinoline, obtained by treating 5-chloro-2-methyl-4-quinolone

with POCl₃ (cf. De, D., Byers L.D., Krogstad, D.J. *J. Heterocyclic Chem.* **1997**, *34*, 315;

25 ¹H NMR (CDCl₃), δ 2.68 (s, 3H), 7.41 (s, 1H), 7.55 (t, 1H), 7.59 (dd, 1H), 7.95 (dd, 1H)), (125 mg, 0.59 mmol) and 1-*tert*-butoxycarbonylpiperazine (559 mg, 3.0 mmol) are reacted according to method A. The Boc group is cleaved with trifluoroacetic acid (3 mL) in CH₂Cl₂ (5 mL) giving 0.18 g of the piperazine.

¹H NMR (CDCl₃), δ 2.63 (s, 3H), 2.73 (t, 2H), 3.04 (d, 2H), 3.23 (t, 2H), 3.37 (d, 2H),

30 6.78 (s, 1H), 7.40 (dd, 1H), 7.44 (t, 1H), 7.84 (dd, 1H).

Preparation of 5-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-

methylquinoline

5-Chloro-2-methyl-4-(piperazin-1-yl)quinoline (0.18 g, 0.7 mmol), 4-fluorophenyl isocyanate (91 μ L, 0.8 mmol), and DMAP (5 mg) in CH_2Cl_2 (10 mL) are reacted

5 according to method C yielding the product as a colorless solid after column chromatography with hexane-EtOAc.

^1H NMR (CDCl_3), δ 2.66 (s, 3H), 2.80 (m, 2H), 3.50 (m, 4H), 4.05 (m, 2H), 6.45 (br. s, 1H), 6.80 (s, 1H), 6.98 (t, 2H), 7.30 (dd, 2H), 7.46 (dd, 1H), 7.49 (d, 1H), 7.90 (dd, 1H).

10 **Example 5****Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-8-methylquinoline***Preparation of 7-Chloro-8-methyl-4-(piperazin-1-yl)quinoline*

15 4,7-Dichloro-8-methylquinoline [obtained from 3-chloro-2-methylaniline following the protocol of Krogstad et. al. (De, D., Byers L.D., Krogstad, D.J. *J. Heterocyclic Chem.* 1997, 34, 315; ^1H NMR (CDCl_3), δ 2.88 (s, 3H), 7.47 (d, 1H), 7.60 (d, 1H), 8.03 (d, 1H), 8.79 (d, 1H))] (212 mg, 1.0 mmol) and 1-*tert*-butoxycarbonylpiperazine (931 mg, 5.0 mmol) are reacted according to method A. The Boc group is cleaved with
20 trifluoroacetic acid (5 mL) in CH_2Cl_2 (5 mL) giving 80 mg of the piperazine.

Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-8-methylquinoline

25 7-Chloro-8-methyl-4-(piperazin-1-yl)quinoline (80 mg, 0.31 mmol), 4-fluorophenyl isocyanate (57 μ L, 0.5 mmol), and DMAP (2 mg) in THF (10 mL) are reacted according to method C yielding the product as a yellow solid after column chromatography with hexane-EtOAc.
30 ^1H NMR ([D]₆-DMSO), δ 2.71 (s, 3H), 3.65 (br., 4H), 3.75 (m, 4H), 7.07 (t, 2H), 7.15 (d, 1H), 7.46 (dd, 1H), 7.65 (d, 1H), 8.02 (d, 1H), 8.63 (d, 1H), 8.68 (br. s, 1H).

Example 6

Preparation f 7-Chloro-4-[4-(4-fluor phenylaminocarbonyl)piperazin-1-yl]-3-methylquinoline

4,7-Dichloro-3-methylquinoline (De, D., Byers L.D., Krogstad, D.J. *J. Heterocyclic Chem.* 1997, 34, 315) (0.30 g, 1.4 mmol) and 1-*tert*-butoxycarbonylpiperazine (0.78 g, 4.2 mmol) are reacted according to method A though the reaction time is extended to 24 h. The Boc group is cleaved with trifluoroacetic acid (3 mL) in CH₂Cl₂ (3 mL) giving 0.28 g of a yellow oil, which is treated with 4-fluorophenyl isocyanate (125 µL, 1.1 mmol) in THF (10 mL) according to method C yielding the product as a colorless solid after column chromatography with hexane-EtOAc.

¹H NMR ([D]₆-DMSO), δ 2.45 (s, 3H), 3.30 (m, 8H), 7.06 (t, 2H), 7.48 (dd, 2H), 7.55 (dd, 1H), 7.96 (d, 1H), 8.23 (d, 1H), 8.63 (s, 1H), 8.65 (br. s, 1H).

Example 7

15 Preparation of 4-[4-(4-Fluorophenylaminocarbonyl)piperazin-1-yl]-7-methoxy-2-methylquinoline

Preparation of 7-Methoxy-8-methyl-4-(piperazin-1-yl)quinoline

4-Chloro-7-methoxy-8-methylquinoline [obtained from *m*-anisidine following the protocol of Krogstad et. al. (De, D., Byers L.D., Krogstad, D.J. *J. Heterocyclic Chem.* 1997, 34, 315; ¹H NMR (CDCl₃), δ 2.70 (s, 3H), 3.93 (s, 3H), 7.20 (dd, 1H), 7.24 (s, 1H), 7.36 (d, 1H), 8.04 (d, 1H))] (0.31 g, 1.5 mmol) and 1-*tert*-butoxycarbonylpiperazine (1.49 g, 8.0 mmol) are reacted according to method A (reaction time: 3.5 h). The Boc group is cleaved with trifluoroacetic acid (5 mL) in CH₂Cl₂ (5 mL) giving 0.20 g of the piperazine as brown oil.

¹H NMR (CDCl₃), δ 2.61 (s, 3H), 3.13 (m, 4H), 3.17 (m, 4H), 3.87 (s, 3H), 6.59 (s, 1H), 7.02 (dd, 1H), 7.30 (d, 1H), 7.79 (d, 1H).

Preparation of 4-[4-(4-Fluorophenylaminocarbonyl)piperazin-1-yl]-7-methoxy-2-methylquinoline

7-Methoxy-8-methyl-4-(piperazin-1-yl)quinoline (0.20 g, 0.8 mmol) and 4-fluorophenyl

isocyanate (102 μ L, 0.9 mmol) in THF (10 mL) are reacted according to method C yielding the product as a yellow solid after column chromatography with hexane-EtOAc.

¹H NMR ([D]₆-DMSO), δ 2.53 (s, 3H), 3.11 (m, 4H), 3.71 (m, 4H), 3.85 (s, 3H), 6.80 (s, 1H), 7.06 (t, 2H), 7.09 (dd, 1H), 7.24 (d, 1H), 7.46 (m, 2H), 7.88 (d, 1H), 8.67 (br. s, 1H).

Example 8

Preparation of 7-Chloro-6-fluoro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-

yl]quinoline

Preparation of 7-Chloro-6-fluoro-4-(piperazin-1-yl)quinoline

4,7-Dichloro-6-fluoroquinoline [obtained from 3-chloro-4-fluoroaniline following the protocol of Krogstad et. al. (De, D., Byers L.D., Krogstad, D.J. *J. Heterocyclic Chem.*

1997, 34, 315; ¹H NMR (CDCl₃), δ 7.51 (d, 1H), 7.94 (d, 1H), 8.22 (d, 1H), 8.76 (d, 1H)]) (0.32 g, 1.7 mmol) and 1-*tert*-butoxycarbonylpiperazine (1.40 g, 7.5 mmol) are reacted according to method A. The Boc group is cleaved with trifluoroacetic acid (5 mL) in CH₂Cl₂ (5 mL) giving 0.36 g of the piperazine as a pale yellow solid.

¹H NMR (CDCl₃), δ 3.15 (m, 8H), 6.86 (d, 1H), 7.70 (d, 1H), 8.11 (d, 1H), 8.69 (d, 1H).

20

Preparation of 7-Chloro-6-fluoro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline

7-Chloro-6-fluoro-4-(piperazin-1-yl)quinoline (0.36 g, 1.35 mmol) and 4-fluorophenyl

25 isocyanate (171 μ L, 1.5 mmol) in THF (10 mL) are reacted according to method C yielding the product as a colorless solid after column chromatography with hexane-EtOAc.

¹H NMR ([D]₆-DMSO) δ 3.17 (m, 4H), 3.71 (m, 4H), 7.07 (t, 2H), 7.10 (d, 1H), 7.46 (dd, 2H), 7.92 (d, 1H), 8.20 (d, 1H), 8.68 (br. s, 1H), 8.72 (d, 1H).

30

Example 9

Preparation of 7-Fluoro-4-[4-(4-fluorophenylaminocarbonyl)pip razin-1-

yl]quinoline*Preparation of 7-Fluoro-4-(piperazin-1-yl)quinoline*

4-Chloro-7-fluoroquinoline [obtained from 3-fluoroaniline following the protocol of
 5 Krogstad et. al. (De, D., Byers L.D., Krogstad, D.J. *J. Heterocyclic Chem.* **1997**, 34,
 315; ^1H NMR (CDCl_3), δ 7.43 (ddd, 1H), 7.46 (d, 1H), 7.75 (dd, 1H), 8.24 (dd, 1H),
 8.77 (d, 1H)]) (0.47 g, 2.6 mmol) and 1-*tert*-butoxycarbonylpiperazine (1.45 g, 7.8
 mmol) are reacted according to method A. The Boc group is cleaved with
 10 trifluoroacetic acid (5 mL) in CH_2Cl_2 (5 mL) giving 0.30 g of the piperazine as yellow
 oil.
 ^1H NMR (CDCl_3) δ 3.14 (m, 4H), 3.19 (m, 4H), 6.80 (d, 1H), 7.24 (ddd, 1H), 7.64 (dd,
 1H), 8.02 (dd, 1H), 8.71 (d, 1H).

Preparation of 7-Fluoro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline

15 7-Fluoro-4-(piperazin-1-yl)quinoline (0.29 g, 1.25 mmol) and 4-fluorophenyl isocyanate
 (142 μL , 1.25 mmol) in THF (10 mL) are reacted according to method C yielding the
 product as colorless crystals after column chromatography with hexane-EtOAc.
 ^1H NMR ([D]₆-DMSO) δ 3.18 (m, 4H), 3.71 (m, 4H), 7.01 (d, 1H), 7.07 (t, 2H), 7.46 (m,
 20 3H), 7.66 (dd, 1H), 8.14 (dd, 1H), 8.68 (br. s, 1H), 8.70 (d, 1H).

Example 10**Preparation of 6,7-Difluoro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline**

25 4-Chloro-6,7-difluoroquinoline [obtained from 3,4-difluoroaniline following the protocol
 of Krogstad et. al. (De, D., Byers L.D., Krogstad, D.J. *J. Heterocyclic Chem.* **1997**, 34,
 315; ^1H NMR (CDCl_3) δ 7.50 (d, 1H), 7.88 (dd, 1H), 7.97 (dd, 1H), 8.75 (d, 1H)]) (0.32
 g, 1.6 mmol) and 1-*tert*-butoxycarbonylpiperazine (0.89 g, 4.8 mmol) are reacted
 30 according to method A. The Boc group is cleaved with trifluoroacetic acid (5 mL) in
 CH_2Cl_2 (5 mL) giving 0.16 g of a yellow solid, which is treated with 4-fluorophenyl
 isocyanate (74 μL , 0.65 mmol) in THF (10 mL) according to method C. The title

product is obtained as colorless needles after column chromatography with hexane-EtOAc.

¹H NMR ([D]₆-DMSO) δ 3.17 (m, 4H), 3.71 (m, 4H), 7.06 (t, 2H), 7.08 (d, 1H), 7.46 (dd, 2H), 7.94 (m, 2H), 8.68 (br. s, 1H), 8.71 (d, 1H).

5

Example 11

Preparation of 7-Cyano-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline

10 4-Chloro-7-cyanoquinoline [obtained from 3-aminobenzonitrile following the protocol of Krogstad et. al. (De, D., Byers L.D., Krogstad, D.J. *J. Heterocyclic Chem.* **1997**, *34*, 315; ¹H NMR (CDCl₃), δ 7.63 (d, 1H), 7.79 (dd, 1H), 8.35 (d, 1H), 8.49 (d, 1H), 8.90 (d, 1H))] (0.29 g, 1.5 mmol) and 1-*tert*-butoxycarbonylpiperazine (0.56 g, 3.0 mmol) are reacted according to method A. The Boc group is cleaved with trifluoroacetic acid (3 mL) in CH₂Cl₂ (5 mL) giving 0.17 g an orange solid, which is treated with 4-fluorophenyl isocyanate (80 μL, 0.67 mmol) in THF (10 mL) according to method C. The title product is obtained as a yellow solid after column chromatography with hexane-EtOAc.

15 ¹H NMR ([D]₆-DMSO) δ 3.20 (m, 4H), 3.73 (m, 4H), 7.06 (t, 2H), 7.16 (d, 1H), 7.46 (dd, 2H), 7.83 (dd, 1H), 8.22 (d, 1H), 8.49 (d, 1H), 8.69 (br. s, 1H), 8.82 (d, 1H).

20

Example 12

Preparation of 4-[4-(4-Fluorophenylaminocarbonyl)piperazin-1-yl]-6,7-methylenedioxyquinoline

25 4-Chloro-6,7-methylenedioxyquinoline [obtained from 3,4-methylendioxyaniline following the protocol of Krogstad et. al. (De, D., Byers L.D., Krogstad, D.J. *J. Heterocyclic Chem.* **1997**, *34*, 315; ¹H NMR ([D]₆-DMSO), δ 6.28 (s, 2H), 7.45 (s, 1H), 7.50 (s, 1H), 7.65 (d, 1H), 8.62 (d, 1H)) (0.50 g, 2.4 mmol) and 1-*tert*-butoxycarbonylpiperazine (1.8 g, 9.6 mmol) are reacted according to method A yielding 391 mg of 4-[4-(*tert*-butoxycarbonyl)piperazin-1-yl]-6,7-methylenedioxyquinoline. After cleaving the Boc group with trifluoroacetic acid (1.5 mL) in CH₂Cl₂ (1.5

mL), the piperazine is treated with 4-fluorophenylisocyanate (86 µL) and triethylamine (0.245 mL) according to method C yielding the title product.

¹H NMR (CDCl₃) δ 3.20 (m, 4H), 3.80 (m, 4H), 6.10 (s, 2H), 6.45 (s, 1H), 6.80 (d, 1H), 7.05 (t, 2H), 7.35 (m, 3H), 7.40 (s, 1H), 8.60 (d, 1H).

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Example 13

Preparation of 4-[4-(4-Fluorophenylaminocarbonyl)piperazin-1-yl]-7-methylquinoline trifluoroacetate

10 4-Chloro-7-methylquinoline [obtained from 3-toluidine following the protocol of Krogstad et. al. (De, D., Byers L.D., Krogstad, D.J. *J. Heterocyclic Chem.* **1997**, *34*, 315; ¹H NMR ([D]₆-DMSO), δ 2.55 (s, 3H), 7.60 (m, 1H), 7.75 (d, 1H), 7.90 (s, 1H), 8.10 (d, 1H), 8.80 (d, 1H)) (0.80 g, 4.5 mmol) and 1-*tert*-butoxycarbonylpiperazine (3.35 g, 18 mmol) are reacted according to method A yielding 475 mg of 4-[4-(*tert*-butoxycarbonyl)piperazin-1-yl]-7-methylquinoline. After cleaving the Boc group with trifluoroacetic acid (1.8 mL) in CH₂Cl₂ (1.8 mL), the piperazine is treated with 4-fluorophenyl isocyanate and triethylamine (0.245 mL) according to method C yielding the title product.

15 ¹H NMR ([D]₆-DMSO) δ 2.53 (s, 3H), 3.60-3.95 (m, 8H), 7.05-7.10 (m, 3H), 7.45-7.50 (m, 3H), 7.70 (s, 1H), 8.12 (d, 1H), 8.60 (d, 1H), 8.65 (s, 1H).

Example 14

Preparation of 4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-7-nitroquinoline

20 4-Chloro-7-nitroquinoline (Ellis, J.; Gellert, E.; Robson, J. *Aust. J. Chem.* **1973**, *26*, 907) (1.8 g, 8.7 mmol) and 1-*tert*-butoxycarbonylpiperazine (5.0 g, 27 mmol) are reacted according to method A furnishing 0.36 g of the piperazine as brown oil after column chromatography with CH₂Cl₂-MeOH.

25 ¹H NMR (CDCl₃) δ 1.5 (s, 9H), 3.2 (m, 4H), 3.8 (m, 4H), 7.0 (d, 1H), 8.2 (d, 1H), 8.25 (dd, 1H), 8.9 (d, 1H), 8.95 (d, 1H).

30 **Example 15**

Preparation of 4-[4-(4-Fluorophenylaminocarbonyl)piperazin-1-yl]-7-nitroquinolin trifluoroacetate

To a solution of 4-[4-(*tert*-butoxycarbonyl)piperazin-1-yl]-7-nitroquinoline (360 mg, 1 mmol) in CH₂Cl₂ (1 mL) is added trifluoroacetic acid (1 mL). After 1½ h at room temp., the mixture is concentrated, and the residue is dissolved in CH₂Cl₂ (20 mL) and 5 treated with triethylamine (0.56 mL, 4 mmol) and 4-fluorophenyl isocyanate (0.16 mL, 1.4 mmol). After 1½ h stirring at room temp., the reaction mixture is washed with water and brine, dried (Na₂SO₄), and concentrated. The residue is purified by reversed phase HPLC with water-MeCN-TFA yielding the title product.

10 ¹H NMR ([D]₆-DMSO) δ 3.65 (m, 4H), 3.75 (m, 4H), 7.1 (m, 2H), 7.25 (d, 1H), 7.5 (m, 2H), 8.3 (m, 1H), 8.4 (d, 1H), 8.7 (s, 1H), 8.75 (m, 1H), 8.8 (d, 1H).

Example 16

Preparation of 7-Amino-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline

15 A suspension of 4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-7-nitroquinoline trifluoroacetate (130 mg) and 10% Pd/C (20 mg) in MeOH is stirred in a hydrogen atmosphere for 30 min. The reaction mixture is filtered and concentrated giving the title product.

20 ¹H NMR (CD₃OD) δ 3.8 (m, 8H), 6.8 (d, 1H), 6.85 (d, 1H), 7.0 (m, 3H), 7.4 (m, 2H), 7.9 (d, 1H), 8.2 (d, 1H).

Example 17

Preparation of 7-Acetylamino-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline, trifluoroacetate

25 A solution of 7-amino-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline (85 mg, 0.23 mmol) and triethylamine (97 μL, 0.70 mmol) in CH₂Cl₂ (20 mL) is treated with acetyl chloride (0.022 mL, 0.28 mmol) and cat. DMAP and stirred at room temp. for 2 h. The reaction mixture is washed with water and brine, dried (Na₂SO₄), and 30 concentrated. The residue is purified by reversed phase HPLC with water-MeCN-TFA yielding the title product.

¹H NMR (CD₃OD) δ 2.2 (s, 3H), 3.8 (m, 4H), 4.0 (m, 4H), 7.0 (m, 2H), 7.1 (d, 1H), 7.4 (m, 2H), 7.6 (m, 1H), 8.2 (d, 1H), 8.4 (d, 1H), 8.6 (s, 1H).

Example 18

5 **Preparation of Ethyl 4-[4-(*tert*-butoxycarbonyl)piperazin-1-yl]-7-chloroquinoline-2-carboxylate**

Preparation of *Ethyl 7-chloro-4-quinolone-2-carboxylate*

3-Chloroaniline (28.5 g, 200 mmol) and diethyl acetylenedicarboxylate (37.4 g, 220 mmol) are heated in MeOH (300 mL) for 3 h at reflux. The reaction mixture is concentrated to yield a yellow residue, which is heated in phenyl ether (400 mL), for ¾ h. The mixture is cooled to room temp. and diluted with hexane whereupon a white solid precipitates which is filtered off: 34.4 g.

10 ¹H NMR (CDCl₃) δ 1.4 (t, 3H), 4.4 (q, 2H), 6.6 (s, 1H), 7.4 (dd, 1H), 8.0 (d, 1H), 8.05 (d, 1H).

Preparation of *Ethyl 4-[4-(*tert*-butoxycarbonyl)piperazin-1-yl]-7-chloroquinoline-2-carboxylate*

Ethyl 7-chloro-4-quinolone-2-carboxylate (1.0 g, 4 mmol) is stirred with sodium hydride (0.144 g, 6 mmol) in DMF (50 mL) for 1 h at 40 °C. *N*-Phenyl (trifluoromethylsulfon)imide (2.0 g, 5.6 mmol) is added and, after an hour, 1-*tert*-butoxycarbonylpiperazine (3.0 g, 6 mmol). The reaction mixture is stirred at 50 °C for 15 h, concentrated, diluted with EtOAc, washed with water and brine, dried (Na₂SO₄), and concentrated. The residue is purified by column chromatography with CH₂Cl₂-MeOH yielding the title product as a colorless solid.

15 ¹H NMR (CDCl₃) δ 1.5 (t, 3H), 1.55 (s, 9H), 3.2 (m, 4H), 3.8 (m, 4H), 4.6 (q, 2H), 7.5 (dd, 1H), 7.65 (s, 1H), 7.95 (d, 1H), 8.25 (d, 1H).

Example 19

20 **Preparation of Ethyl 7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline-2-carboxylate**

To a solution of ethyl 4-[4-(*tert*-butoxycarbonyl)piperazin-1-yl]-7-chloroquinoline-2-carboxylate (1.0 g, 2.4 mmol) in CH₂Cl₂ (5 mL) is added trifluoroacetic acid (5 mL). After 1 h at room temp., the reaction mixture is adjusted to pH 10-11 with 10% aq. NaOH and extracted with EtOAc. The combined extracts are dried (Na₂SO₄) and concentrated. The residue is dissolved in CH₂Cl₂ (70 mL) and treated with 4-fluorophenyl isocyanate (0.32 mL, 2.8 mmol) for 15 h at room temp. The reaction mixture is washed with water and brine, dried (Na₂SO₄), and concentrated. The residue is purified by column chromatography with CH₂Cl₂-MeOH yielding the title product.

10 ¹H NMR (CDCl₃) δ 1.5 (t, 3H), 3.4 (m, 4H), 3.8 (m, 4H), 4.6 (q, 2H), 6.4 (s, 1H), 7.0 (m, 2H), 7.35 (m, 2H), 7.55 (m, 1H), 7.7 (s, 1H), 8.0 (d, 1H), 8.3 (d, 1H).

Example 20

Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-(hydroxymethyl)quinoline, trifluoroacetate

Ethyl 7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline-2-carboxylate (380 mg, 0.83 mmol) and a 1 M THF solution of super hydride (6.7 mL, 6.7 mmol) are stirred at room temp. for $\frac{2}{3}$ h. Water (10 mL) and 10% aq. NaOH (10 mL) is added, and stirring is continued for $\frac{2}{3}$ h. The mixture is acidified with 5 N HCl (pH 5) and extracted with EtOAc. The extracts are dried (Na₂SO₄) and concentrated. The residue is purified by reversed phase HPLC with water-MeCN-TFA yielding the title product.

25 ¹H NMR ([D]₆-DMSO) δ 3.7 (m, 4H), 3.8 (m, 4H), 4.8 (s, 2H), 7.05 (m, 2H), 7.15 (s, 1H), 7.45 (m, 2H), 7.65 (m, 1H), 8.7 (m, 2H), 8.65 (s, 1H).

Example 21

Preparation of 2-Azidomethyl-7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline

30 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-(hydroxymethyl)quinoline, trifluoroacetate (100 mg, 0.24 mmol) diphenylphosphoryl azide (DPPA, 0.052 mL, 0.24 mmol), and DBU (0.036 mL, 0.24 mmol) are stirred in

THF (30 mL) for 2 h at room temp. The reaction mixture is diluted with EtOAc, washed with water and brine, dried (Na_2SO_4) and concentrated. The residue is purified by column chromatography with CH_2Cl_2 -MeOH to give the title product as a yellow oil.

^1H NMR (CDCl_3) δ 3.3 (m, 4H), 3.8 (m, 4H), 4.6 (s, 2H), 6.4 (s, 1H), 7.0 (m, 2H), 7.3

5 (m, 2H), 7.5 (dd, 1H), 7.9 (d, 1H), 8.0 (s, 1H).

Example 22

Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-(methoxymethyl)quinoline, trifluoroacetate

10

and

Example 23

Preparation of 2-Aminomethyl-7-chloro-4-[4-(4-

15

fluorophenylaminocarbonyl)piperazin-1-yl]quinoline, trifluoroacetate

A solution of 2-azidomethyl-7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline (24 mg, 0.055 mmol) in MeOH (5 mL) is added to a suspension of tin(II) chloride (32 mg, 0.055 mmol) in MeOH (10 mL). After stirring for 2 h at room temp. the 20 reaction mixture is concentrated, and the residue is suspended in water. The mixture is adjusted with 10% aq. NaOH (pH 11-12) and extracted with EtOAc. The combined extracts are dried (Na_2SO_4) and concentrated. The residue is purified by reversed phase HPLC with water-MeCN-TFA yielding 7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-(methoxymethyl)quinoline, trifluoroacetate
25 and 2-aminomethyl-7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline, trifluoroacetate.

7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-(methoxymethyl)quinoline, trifluoroacetate

30 ^1H NMR (CD_3OD) δ 3.6 (s, 3H), 3.9 (m, 4H), 4.0 (m, 4H), 4.8 (s, 2H), 7.0 (m, 2H), 7.2 (s, 1H), 7.4 (m, 2H), 7.7 (m, 1H), 8.1 (d, 1H), 8.3 (d, 1H).

2-Aminomethyl-7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline,

trifluoroacetate

¹H NMR (CD₃OD) δ 3.6 (m, 4H), 3.8 (m, 4H), 4.4 (s, 2H), 7.0 (m, 2H), 7.1 (s, 1H), 7.3 (m, 2H), 7.6 (m, 1H), 8.0 (d, 1H), 8.2 (d, 1H).

5 **Example 24**

Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline-2-carboxylic acid

A solution of ethyl 7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline-2-carboxylate (389 mg, 0.85 mmol) in THF-water 3:1 (5 mL) is treated with LiOH x H₂O (126 mg, 3 mmol) and stirred at room temp. for 4 h. The reaction mixture is concentrated and acidified with 1 N HCl (pH 4). The colorless precipitate is collected by filtration yielding the title product.

¹H NMR (CD₃OD) δ 3.50 (m, 4H), 3.75 (m, 4H), 7.00 (t, 2H), 7.40 (m, 2H), 7.55 (s, 1H), 7.60 (m, 1H), 8.20 (m, 2H).

Example 25

Preparation of 2-*tert*-Butoxycarbonylamino-7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline

20 A solution of 7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline-2-carboxylic acid (215 mg, 0.5 mmol), diphenylphosphoryl azide (DPPA, 144 μL, 0.65 mmol), and triethylamine (92 μL, 0.65 mmol) in *t*-butanol (5 mL) is heated at 105 °C in an Ar atmosphere for 3 h. The solvent is removed, and the residue is dissolved in EtOAc, washed with water, dried (MgSO₄), and concentrated. Purification by column chromatography with CH₂Cl₂-MeOH affords the title product.

¹H NMR ([D]₆-DMSO) δ 1.35 (s, 9H), 3.05 (m, 4H), 3.60 (m, 4H), 6.95 (t, 2H), 7.10 (m, 1H), 7.20-7.40 (m, 3H), 7.55 (m, 2H), 7.85 (d, 1H), 8.60 (s, 1H).

30 **Example 26**

Preparation of 2-Amino-7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline, trifluoroacetate

2-*tert*-Butoxycarbonylamino-7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline (440 mg, 0.88 mmol) in CH₂Cl₂ (1.5 mL) is treated with trifluoroacetic acid (1.5 mL) for 2 h at room temp. The solvent is removed, and the residue purified by

5 reversed phase HPLC with water-MeCN-TFA giving the title product.

¹H NMR (CD₃OD) δ 3.45 (m, 4H), 3.80 (m, 4H), 6.35 (s, 1H), 7.00 (t, 2H), 7.36 (m, 2H), 7.45 (d, 1H), 7.60 (s, 1H), 8.00 (d, 1H).

Example 27

10 **Preparation of 2-Acetylamino-7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline**

2-*tert*-Butoxycarbonylamino-7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline (110 mg, 0.22 mmol) in CH₂Cl₂ (1 mL) is treated with trifluoroacetic acid (1 mL) for 2 h at room temp. The reaction mixture is concentrated. The residue is dissolved in CH₂Cl₂ (9 mL). Triethylamine (154 μL, 1.1 mmol), acetyl chloride (26 μL, 0.24 mmol), and DMAP (10 mg) are added. The reaction mixture is stirred at room temp. for 3 h, washed with water, dried (MgSO₄), and concentrated. Purification of the residue by column chromatography with CH₂Cl₂-MeOH affords the title product.

15 20 ¹H NMR (CDCl₃) δ 2.25 (s, 3H), 3.30 (m, 4H), 3.80 (m, 4H), 7.00 (t, 2H), 7.35 (m, 4H), 7.75 (s, 1H), 7.90 (d, 1H), 7.95 (s, 1H).

Example 28

25 **Preparation of N-[7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinolin-2-yl]urea, trifluoroacetate**

To a solution of 2-amino-7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline (42 mg, 0.11 mmol) in 1,4-dioxane-THF 3:1 (2 mL) is added trimethylsilyl isocyanate (0.1 mL). The reaction mixture is heated in a sealed tube for 18 h at 80 °C.

30 More isocyanate (0.2 mL) is added, and the mixture is heated for further 18 h. The solvent is removed, and the residue is purified by flash chromatography with CH₂Cl₂-MeOH followed by reversed phase HPLC with water-MeCN-TFA affording the title

product. ^1H NMR (CD_3OD) δ 3.65 (m, 4H), 3.80 (m, 4H), 6.55 (s, 1H), 7.00 (t, 2H), 7.38 (m, 2H), 7.60 (d, 1H), 8.00 (s, 1H), 8.10 (d, 1H).

Example 29

5 **Preparation of N-[7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinolin-2-yl]-N'-phenylurea**

To a solution of 2-amino-7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline (20 mg, 0.05 mmol) in MeCN-1,2-dichloroethane 1:1 (12 mL) is added phenyl isocyanate (17 μL , 0.11 mmol). The reaction mixture is heated at reflux for 1 h.

10 Upon cooling the title product precipitates and is collected by filtration.

^1H NMR ($[\text{D}]_6\text{-DMSO}$) δ 3.16 (m, 4H), 3.70 (m, 4H), 6.90 (s, 1H), 7.05 (m, 3H), 7.35 (t, 2H), 7.40 (m, 1H), 7.45 (m, 2H), 7.65 (d, 2H), 7.95 (m, 2H).

15 **Example 30**

Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-(methylamino)quinoline, trifluoroacetate

To a solution of 2-*tert*-butoxycarbonylamino-4-[4-*tert*-butoxycarbonylpiperazin-1-yl]-7-chloroquinoline (150 mg, 0.35 mmol) in THF (2 mL) is added a suspension of sodium hydride (25 mg, 1.1 mmol) in THF (4 mL). After 30 min at room temp. iodomethane (33 μL , 0.53 mmol) is added, and stirring is continued for 7 h. The reaction mixture is diluted with EtOAc, washed with water, dried (MgSO_4), and concentrated. The residue is purified by column chromatography with $\text{CH}_2\text{Cl}_2\text{-MeOH}$ giving 86 mg of 2-(*N*-*t*-butoxycarbonyl-*N*-methylamino)-4-[4-*tert*-butoxycarbonylpiperazin-1-yl]-7-chloroquinoline. This is treated with TFA- CH_2Cl_2 1:1 (1mL) to cleave the Boc groups and 4-fluorophenyl isocyanate (30 μL) and triethylamine (0.20 mL) according to method C affording the title product.

^1H NMR (CD_3OD) δ 3.10 (s, 3H), 3.40 (m, 4H), 3.80 (m, 4H), 6.35 (s, 1H), 7.00 (t, 2H), 7.35 (m, 2H), 7.45 (dd, 1H), 7.80 (m, 1H), 8.00 (d, 1H).

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Example 31

Preparation of 4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-7-chloro-2-quinolone

Preparation of 4,7-Dichloro-2-quinolone

4,7-Dichloroquinoline (20.4 g) and peracetic acid (32 wt.% [from Aldrich], 24 mL) are heated at reflux for 8 h and kept at room temp. for 18 h. The precipitate is collected and washed with hexane to give 23 g of 4,7-dichloroquinoline-N-oxide. 1.0 g (4.7 mmol) thereof is heated in phosphoryl chloride (3.7 mL) for 1½ h at reflux. The reaction mixture is concentrated and the residue is treated with 10% aq. NaOH (pH 9). The precipitate is collected, washed with water, and dried yielding 1.0 g of 2,4,7-trichloroquinoline. 250 mg thereof is heated with 15% H₂SO₄ (20 mL) and 1,4-dioxane (10 mL) in a sealed tube at 140 °C for 8 h. The mixture is kept at room temp. for 18 h. The precipitate is collected, washed with water, and dried affording 195 mg of 4,7-dichloro-2-quinolone.

¹H NMR (CDCl₃) δ 7.55 (s, 1H), 7.60 (d, 1H), 8.03 (s, 1H), 8.16 (d, 1H).

Preparation of 4-[4-(tert-Butoxycarbonyl)piperazin-1-yl]-7-chloro-2-quinolone
 4,7-Dichloro-2-quinolone (192 mg, 0.88 mmol) and 1-tert-butoxycarbonylpiperazine (580 mg, 3.1 mmol) in *n*-butanol (10 mL) are heated in a sealed tube for 2 h at 150 °C, when additional piperazine (200 mg) is added and heating is continued for further 18 h. After removing the solvent, the residue is purified by column chromatography with CH₂Cl₂-MeOH-NH₄OH yielding the title product as a colorless solid.
¹H NMR ([D]₆-DMSO) δ 1.4 (s, 9H), 3.0 (m, 4H), 3.5 (m, 4H), 5.9 (s, 1H), 7.15 (d, 1H), 7.3 (s, 1H), 7.7 (d, 1H).

Example 32**Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-quinolone**

4-[4-(tert-Butoxycarbonyl)piperazin-1-yl]-7-chloro-2-quinolone (100 mg) is deprotected with TFA-CH₂Cl₂ 1:1 (0.5 mL) and transformed to the title product with 4-fluorophenyl isocyanate (50 µL) according to method C.

¹H NMR ([D]₆-DMSO) δ 3.15 (m, 4H), 3.65 (m, 4H), 5.90 (s, 1H), 7.05 (t, 2H), 7.18 (d, 1H), 7.30 (s, 1H), 7.45 (m, 2H), 7.70 (d, 1H), 8.65 (s, 1H), 11.40 (s, 1H).

Example 33**Preparation of 7-Chloro-2-dimethylamino-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline**

5

4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-7-chloroquinol-2-one (74 mg, 0.2 mmol), sodium hydride (8 mg, 0.3 mmol), *N*-phenyl(trifluoromethylsulfon)imide (101 mg, 0.28 mmol), and dimethylamine (2 M in THF, 2 mL, 4.0 mmol) are treated according to method E yielding 46 mg of the 4-[4-(*tert*-butoxycarbonyl)piperazin-1-yl]-7-chloro-2-dimethylaminoquinoline. This is treated with TFA-CH₂Cl₂ 1:1 (0.6 mL) for 1 h and concentrated. The residue is converted into the title product with 4-fluorophenyl isocyanate (19 µL) and triethylamine (0.11 mL).

10 ¹H NMR (CDCl₃) δ 3.20 (m, 10H), 3.75 (m, 4H), 6.30 (s, 1H), 6.42 (s, 1H), 7.00 (t, 2H), 7.05 (d, 1H), 7.35 (m, 2H), 7.75 (m, 2H).

15

Example 34**Preparation of 7-Chloro-2-ethylamino-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline**

20 To a solution of 2-*tert*-butoxycarbonylamino-4-[4-*tert*-butoxycarbonylpiperazin-1-yl]-7-chloroquinoline (135 mg, 0.29 mmol) in THF (2 mL) is added a suspension of sodium hydride (21 mg, 0.87 mmol) in THF (4 mL). After 3/4 h at room temp. iodoethane (0.45 mmol) in THF (1 mL) is added, and stirring is continued for 18 h. More iodoethane (0.42 mmol) is added, and stirring is continued at 45 °C for 18 h. The reaction mixture is diluted with EtOAc, washed with water, dried (MgSO₄), and concentrated. The residue is purified by column chromatography with CH₂Cl₂-MeOH giving 60 mg of 2-(*N*-*tert*-butoxycarbonyl-*N*-ethylamino)-4-[4-*tert*-butoxycarbonylpiperazin-1-yl]-7-chloroquinoline. This is treated with TFA-CH₂Cl₂ 1:1 (1mL) to cleave the Boc groups. 4-Fluorophenyl isocyanate (24 µL) and triethylamine (0.14 mL) are added according to method C affording the title product.

30 ¹H NMR (CD₃OD) δ 1.25 (t, 3H), 3.15 (m, 4H), 3.45 (q, 2H), 3.80 (m, 4H), 6.25 (s, 1H), 7.00 (t, 2H), 7.10 (m, 1H), 7.35 (m, 2H), 7.55 (s, 1H), 7.80 (d, 1H).

Example 35**Preparation of 7-Chloro-2-[2-(dimethylamino)ethylamino]-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline**

5

4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-7-chloroquinol-2-one (145 mg, 0.4 mmol), sodium hydride (15 mg, 0.6 mmol), *N*-phenyl (trifluoromethylsulfon)imide (202 mg, 0.56 mmol), and N,N-dimethylethylenediamine (276 µL, 2.4 mmol) are treated according to method E yielding 98 mg of 4-[4-(*tert*-butoxycarbonyl)piperazin-1-yl]-7-chloro-2-[2-(dimethylamino)ethylamino]quinoline. 38 mg thereof are treated with TFA-CH₂Cl₂ 1:1 (0.6 mL) for 1 h and concentrated. The residue is converted into the title product with 4-fluorophenyl isocyanate (16 µL) and triethylamine (120 µL).

¹H NMR ([D]₆-DMSO) δ 2.20 (s, 6H), 2.40 (t, 2H), 3.00 (m, 4H), 3.40 (m, 2H), 3.70 (m, 4H), 6.38 (s, 1H), 6.90 (m, 1H), 7.05 (m, 3H), 7.40 (s, 1H), 7.45 (m, 2H), 7.75 (d, 1H), 15 8.65 (s, 1H).

Example 36**Preparation of 7-Chloro-2-cyclohexylamino-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline**

20

4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-7-chloroquinol-2-one (150 mg, 0.4 mmol), sodium hydride (15 mg, 0.6 mmol), *N*-phenyl (trifluoromethylsulfon)imide (208 mg, 0.56 mmol), and cyclohexylamine (286 µL, 2.4 mmol) are treated according to method E yielding 21 mg of 4-[4-(*tert*-butoxycarbonyl)piperazin-1-yl]-7-chloro-2-(cyclohexylamino)quinoline which is treated with TFA-CH₂Cl₂ 1:1 (0.6 mL) for 1 h and concentrated. The residue is converted into the title product with 4-fluorophenyl isocyanate (8 µL) and triethylamine (60 µL).

¹H NMR (CD₃OD) δ 1.30 (m, 3H), 1.45 (m, 2H), 1.70 (m, 1H), 1.80 (m, 2H), 2.00 (m, 2H), 3.10 (m, 4H), 3.75 (m, 4H), 3.90 (m, 1H), 6.25 (s, 1H), 7.00 (t, 2H), 7.10 (d, 1H), 30 7.38 (m, 2H), 7.60 (s, 1H), 7.80 (d, 1H).

Example 37

Preparation of 7-Chloro-2-cyclopropylamino-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinolin

4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-7-chloroquinol-2-one (150 mg, 0.4 mmol), sodium hydride (15 mg, 0.6 mmol), *N*-phenyl (trifluoromethylsulfon)imide (208 mg, 0.56 mmol), and cyclopropylamine (580 µL, 6.4 mmol) are treated according to method E yielding 75 mg of 4-[4-(*tert*-butoxycarbonyl)piperazin-1-yl]-7-chloro-2-(cyclopropylamino)quinoline. 62 mg thereof are treated with TFA-CH₂Cl₂ 1:1 (0.6 mL) for 1 h and concentrated. The residue is converted into the title product with 4-fluorophenyl isocyanate (26 µL) and triethylamine (193 µL).

¹H NMR (CD₃OD) δ 0.55 (m, 2H), 0.85 (m, 2H), 2.75 (m, 1H), 3.19 (m, 4H), 3.80 (m, 4H), 6.40 (s, 1H), 7.00 (t, 2H), 7.16 (d, 1H), 7.36 (m, 2H), 7.60 (s, 1H), 7.85 (d, 1H).

Example 38

Preparation of 7-Chloro-2-cyclopropylmethylamino-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline

4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-7-chloroquinol-2-one (150 mg, 0.4 mmol), sodium hydride (15 mg, 0.6 mmol), *N*-phenyl (trifluoromethylsulfon)imide (208 mg, 0.56 mmol), and cyclopropylmethylamine (360 µL, 4.0 mmol) are treated according to method E yielding 53 mg of 4-[4-(*tert*-butoxycarbonyl)piperazin-1-yl]-7-chloro-2-(cyclopropylmethylamino)quinoline. 47 mg thereof are treated with TFA-CH₂Cl₂ 1:1 (0.6 mL) for 1 h and concentrated. The residue is converted into the title product with 4-fluorophenyl isocyanate (19 µL) and triethylamine (140 µL).

¹H NMR (CD₃OD), δ 0.30 (m, 2H), 0.55 (m, 2H), 1.10 (m, 1H), 3.10 (m, 4H), 3.20 (m, 2H), 3.80 (m, 4H), 6.30 (s, 1H), 7.00 (t, 2H), 7.15 (d, 1H), 7.40 (m, 2H), 7.55 (s, 1H), 7.80 (d, 1H).

Example 39

Preparation of 4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-7-chloro-2-(2-hydroxyethylamino)quinoline

4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-7-chloroquinol-2-one (150 mg, 0.4 mmol), sodium hydride (15 mg, 0.6 mmol), *N*-phenyl (trifluoromethylsulfon)imide (208 mg, 0.56 mmol), and ethanolamine (100 μ L, 1.65 mmol) are treated according to method E yielding the title product.

5 ^1H NMR (CDCl_3) δ 1.5 (s, 9H), 3.1 (m, 4H), 3.7 (m, 6H), 3.9 (m, 2H), 6.1 (s, 1H), 7.15 (dd, 1H), 7.65 (d, 1H), 7.7 (d, 1H).

Example 40

Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-[2-(4-fluorophenylaminocarbonyloxy)ethylamino]quinoline

4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-7-chloro-2-(2-hydroxyethylamino)quinoline (62 mg, 0.15 mmol) is deprotected with TFA followed by reaction with 4-fluorophenyl isocyanate (24 μ L, 0.21 mmol) and triethylamine (150 μ L, 1.1 mmol) according to
15 method C yielding the title product as a colorless solid after column chromatography with CH_2Cl_2 -MeOH.

^1H NMR (CDCl_3) δ 3.2 (m, 4H), 3.75 (m, 4H), 3.85 (m, 2H), 4.4 (m, 2H), 5.1 (s, 1H), 6.1 (s, 1H), 6.4 (s, 1H), 6.7 (s, 1H), 7.0 (m, 4H), 7.15 (dd, 1H), 7.3 (m, 4H), 7.65 (d, 1H), 7.7 (d, 1H).

20

Example 41

Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-(2-hydroxyethylamino)quinoline trifluoroacetate

25 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-[2-(4-fluorophenylaminocarbonyloxy)ethylamino]quinoline (39 mg, 0.07 mmol) and 6 N KOH (1.1 mL, 6.6 mmol) in EtOH-THF 5:1 (6 mL) are heated at reflux for several hours until the starting material is consumed (TLC). The reaction mixture is diluted with water and extracted with EtOAc. The organic phase is dried (MgSO_4) and concentrated. The
30 residue is purified by reversed phase HPLC with water-MeCN-TFA affording the title product.

^1H NMR (CD_3OD) δ 3.40 (m, 4H), 3.65 (m, 2H), 3.85 (m, 6H), 6.40 (s, 1H), 7.00 (t, 2H),

7.35 (m, 2H), 7.45 (d, 1H), 7.75 (s, 1H), 8.00 (d, 1H).

Example 42

Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-(2-methoxyethyl)aminoquinoline

4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-7-chloroquinol-2-one (150 mg, 0.4 mmol), sodium hydride (15 mg, 0.6 mmol), *N*-phenyl(trifluoromethylsulfon)imide (208 mg, 0.56 mmol), and 2-methoxyethylamine (362 µL, 4.0 mmol) are treated according to method E yielding 85 mg of 4-[4-(*tert*-butoxycarbonyl)piperazin-1-yl]-7-chloro-2-(2-methoxyethylamino)quinoline. 74 mg thereof are treated with TFA-CH₂Cl₂ 1:1 (0.6 mL) for 1 h and concentrated. The residue is converted into the title product with 4-fluorophenyl isocyanate (30 µL) and triethylamine (230 µL).

¹H NMR (CD₃OD) δ 3.15 (m, 4H), 3.40 (s, 3H), 3.60 (m, 6H), 3.80 (m, 4H), 6.35 (s, 1H), 7.00 (t, 2H), 7.10 (d, 1H), 7.35 (m, 2H), 7.55 (s, 1H), 7.80 (d, 1H).

Example 43

Preparation of 4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-7-chloro-2-(*n*-propylamino)quinoline

4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-7-chloroquinol-2-one (150 mg, 0.41 mmol), sodium hydride (15 mg, 0.6 mmol), *N*-phenyl(trifluoromethylsulfon)imide (208 mg, 0.58 mmol), and propylamine (0.14 mL, 1.65 mmol) are treated according to method E yielding the title product.

¹H NMR (CDCl₃) δ 1.0 (t, 3H), 1.5 (s, 9H), 1.7 (m, 2H), 3.1 (m, 4H), 3.4 (q, 2H), 3.7 (m, 4H), 6.1 (s, 1H), 7.1 (dd, 1H), 7.6 (d, 1H), 7.7 (d, 1H).

Example 44

Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-(*n*-propylamino)quinoline

4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-7-chloro-2-(*n*-propylamino)quinoline (25 mg,

0.06 mmol) is deprotected with TFA in CH₂Cl₂ and transformed into the title product according to method C using 4-fluorophenyl isocyanate (10 µL, 0.09 mmol) and triethylamine (60 µL, 0.43 mmol) to give a colorless solid.

10 ¹H NMR (CDCl₃) δ 1.0 (t, 3H), 1.8 (m, 2H), 3.2 (m, 4H), 3.4 (m, 2H), 3.8 (m, 4H), 6.1
5 (s, 1H), 6.4 (s, 1H), 7.0 (m, 2H), 7.1 (m, 1H), 7.3 (m, 2H), 7.65 (d, 1H), 7.7 (d, 1H).

Example 45

Preparation of 4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-7-chloro-2-(isopropylamino)-quinoline

10 4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-7-chloroquinol-2-one (150 mg, 0.41 mmol), sodium hydride (15 mg, 0.6 mmol), *N*-phenyl (trifluoromethylsulfon)imide (208 mg, 0.58 mmol), and isopropylamine (0.14 mL, 1.65 mmol) are treated according to method E yielding the title product.
15 ¹H NMR (CDCl₃) δ 1.25 (d, 6H), 1.5 (s, 9H), 3.1 (m, 4H), 3.7 (m, 4H), 4.2 (m, 1H), 6.0 (s, 1H), 7.1 (dd, 1H), 7.6 (d, 1H), 7.7 (d, 1H).

Example 46

Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-(isopropylamino)quinoline

20 4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-7-chloro-2-(isopropylamino)quinoline (33 mg, 0.08 mmol) is deprotected with TFA in CH₂Cl₂ and transformed into the title product according to method C using 4-fluorophenyl isocyanate (13 µL, 0.11 mmol) and triethylamine (80 µL, 0.57 mmol) to give a colorless solid.
25 ¹H NMR (CDCl₃) δ 1.2 (d, 6H), 3.2 (m, 4H), 3.8 (m, 4H), 4.2 (m, 1H), 4.6 (m, 1H), 6.0 (s, 1H), 6.4 (s, 1H), 7.0 (m, 2H), 7.1 (m, 1H), 7.3 (m, 2H), 7.6 (d, 1H), 7.7 (d, 1H).

Example 47

30 **Preparation of 4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-2-(*n*-butylamino)-7-chloroquinoline**

53001A

4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-7-chloroquinol-2-one (150 mg, 0.41 mmol), sodium hydride (15 mg, 0.6 mmol), *N*-phenyl (trifluoromethylsulfon)imide (208 mg, 0.58 mmol), and *n*-butylamine (0.16 mL, 1.65 mmol) are treated according to method E yielding the title product.

5 ^1H NMR (CDCl_3) δ 1.0 (t, 3H), 1.45 (m, 2H), 1.5 (s, 9H), 3.1 (m, 4H), 3.4 (q, 2H), 3.7 (m, 4H), 6.0 (s, 1H), 7.1 (dd, 1H), 7.6 (d, 1H), 7.7 (d, 1H).

Example 48

Preparation of 2-*n*-Butylamino-7-chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline

4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-2-(*n*-butylamino)-7-chloroquinoline (47 mg, 0.11 mmol) is deprotected with TFA in CH_2Cl_2 and transformed into the title product according to method C using 4-fluorophenyl isocyanate (18 μL , 0.16 mmol) and triethylamine (110 μL , 0.79 mmol) to give a colorless solid.

15 ^1H NMR (CDCl_3) δ 1.0 (t, 3H), 1.5 (m, 2H), 1.6 (m, 2H), 3.2 (m, 4H), 3.4 (m, 2H), 3.8 (m, 4H), 4.6 (m, 1H), 6.1 (s, 1H), 6.4 (s, 1H), 7.0 (m, 2H), 7.1 (m, 1H), 7.3 (m, 2H), 7.65 (d, 1H), 7.7 (d, 1H).

Example 49

Preparation of 4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-7-chloro-2-(*n*-pentylamino)quinoline

4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-7-chloroquinol-2-one (150 mg, 0.41 mmol), sodium hydride (15 mg, 0.6 mmol), *N*-phenyl (trifluoromethylsulfon)imide (208 mg, 0.58 mmol), and *n*-pentylamine (0.19 mL, 1.65 mmol) are treated according to method E yielding the title product.

30 ^1H NMR (CDCl_3) δ 0.9 (t, 3H), 1.4 (m, 4H), 1.5 (s, 9H), 1.65 (m, 2H), 3.1 (m, 4H), 3.4 (q, 2H), 3.7 (m, 4H), 6.1 (s, 1H), 7.1 (dd, 1H), 7.6 (d, 1H), 7.7 (d, 1H).

30

Example 50

Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-(*n*-

pentylamino)quinoline

4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-7-chloro-2-(*n*-pentylamino)quinoline (68 mg, 0.11 mmol) is deprotected with TFA in CH₂Cl₂ and transformed into the title product

5 according to method C using 4-fluorophenyl isocyanate (18 µL, 0.16 mmol) and triethylamine (110 µL, 0.79 mmol) to give a colorless solid.

¹H NMR (CDCl₃) δ 0.9 (t, 3H), 1.4 (m, 4H), 1.7 (m, 2H), 3.2 (m, 4H), 3.4 (m, 2H), 3.8 (m, 4H), 4.6 (m, 1H), 6.0 (s, 1H), 6.4 (s, 1H), 7.0 (m, 2H), 7.1 (m, 1H), 7.3 (m, 2H), 7.65 (d, 1H), 7.7 (d, 1H).

10

Example 51**Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-[2-(4-fluorophenyl)ethylamino]quinoline**

15 4-[4-(*tert*-Butoxycarbonyl)piperazin-1-yl]-7-chloroquinol-2-one (150 mg, 0.41 mmol), sodium hydride (15 mg, 0.6 mmol), *N*-phenyl (trifluoromethylsulfon)imide (208 mg, 0.58 mmol), and 4-fluorophenethylamine (0.22 mL, 1.65 mmol) are treated according to method E yielding 4-[4-(butoxycarbonyl)piperazin-1-yl]-7-chloro-2-[2-(4-fluorophenyl)ethylamino]quinoline, which is deprotected with TFA in CH₂Cl₂ and

20 transformed into the title product according to method C using 4-fluorophenyl isocyanate (18 µL, 0.16 mmol) and triethylamine (110 µL, 0.79 mmol) to give a colorless solid.

¹H NMR (CDCl₃) δ 3.0 (t, 2H), 3.2 (m, 4H), 3.8 (m, 6H), 6.0 (s, 1H), 6.4 (s, 1H), 7.0 (m, 4H), 7.1 (m, 1H), 7.2 (m, 2H), 7.3 (m, 2H), 7.7 (m, 2H).

25

Example 52**Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)-3-methylpiperazin-1-yl]quinoline**

30 *Preparation of 7-Chloro-4-(3-methylpiperazin-1-yl)quinoline*

4,7-Dichloroquinoline (0.5 g, 2.5 mmol) and 2-methylpiperazine (1.25 g, 12.5 mmol) are reacted according to method A yielding the product as a brown oil.

¹H NMR (CDCl₃) δ 1.55 (d, 3H), 2.52 (dd, 1H), 2.88 (td, 1H), 3.19 (m, 3H), 3.42 (m, 2H), 6.82 (d, 1H), 7.40 (dd, 1H), 7.93 (d, 1H), 8.02 (d, 1H), 8.69 (d, 1H).

Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)-3-methylpiperazin-1-

⁵ *yl]quinoline*

7-Chloro-4-(3-methylpiperazin-1-yl)quinoline (0.13 g, 0.5 mmol) and 4-fluorophenyl isocyanate (57 μL, 0.5 mmol) in CH₂Cl₂ (10 mL) are reacted according to method C yielding the title product after column chromatography with hexane-EtOAc.

¹⁰ ¹H NMR ([D]₆-DMSO) δ 1.41 (d, 3H), 2.83 (m, 1H), 2.99 (m, 1H), 3.40 (m, 2H), 3.50 (m, 1H), 4.04 (m, 1H), 4.50 (m, 1H), 7.05 (m, 3H), 7.46 (m, 2H), 7.59 (dd, 1H), 7.98 (d, 1H), 8.11 (d, 1H), 8.60 (br. s, 1H), 8.71 (d, 1H).

Example 53

¹⁵ **Preparation of 4-[4-(*tert*-Butoxycarbonyl)-2-methylpiperazin-1-yl]-7-chloroquinoline**

Sodium hydride (72 mg, 3 mmol) is added to a solution of 7-chloro-4-quinolone (359 mg, 2 mmol) in DMF. After 1 h at 40 °C *N*-phenyl (trifluoromethylsulfon)imide (1.0 g, 2.8 mmol) and, 1 h later, 1-*tert*-butoxycarbonyl-3-methylpiperazine (1.8 g, 8 mmol) is added. The reaction mixture is stirred at 80 °C for 2 days, concentrated, diluted with EtOAc, washed with water and brine, dried (Na₂SO₄), and concentrated. The residue is purified by reversed phase HPLC with water-MeCN-TFA to give the title product.

²⁰ ¹H NMR (CDCl₃) δ 1.0 (d, 3H), 1.5 (s, 9H), 2.9 (m, 1H), 3.3-3.8 (m, 6H), 6.9 (d, 1H), 7.5 (m, 1H), 8.1 (m, 2H), 8.8 (d, 1H).

Example 54

Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)-2-methylpiperazin-1-yl]quinoline

³⁰

4-[4-(*tert*-butoxycarbonyl)-2-methylpiperazin-1-yl]-7-chloroquinoline (0.30 g, 0.8 mmol) is deprotected with TFA in CH₂Cl₂ and transformed after being washed with base into

the product according to method C using 4-fluorophenyl isocyanate (61 μ L, 0.54 mmol) to give the title product a colorless solid.

^1H NMR (CDCl_3) δ 1.0 (d, 3H), 3.0 (m, 1H), 3.5 (m, 2H), 3.6-3.9 (m, 4H), 6.4 (s, 1H), 6.9 (d, 1H), 7.0 (m, 2H), 7.3 (m, 2H), 7.5 (m, 1H), 8.1 (m, 2H), 8.8 (d, 1H).

5

Example 55

Preparation of 7-Chloro-4-[3,5-dimethyl-4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline

10 *Preparation of 7-Chloro-4-(3,5-dimethylpiperazin-1-yl)quinoline*

4,7-Dichloroquinoline (0.40 g, 2.0 mmol) and 2,6-dimethylpiperazine (0.57 g, 5.0 mmol) in *n*-butanol (20 mL) are reacted according to method B yielding the title product after column chromatography with CH_2Cl_2 -MeOH.

15 ^1H NMR (CDCl_3) δ 1.2 (d, 6H), 2.5 (t, 2H), 3.2 (m, 2H), 3.4 (d, 2H), 6.8 (d, 1H), 7.4 (d, 1H), 7.95 (d, 1H), 8.05 (s, 1H), 8.7 (d, 1H).

Preparation of 7-Chloro-4-[3,5-dimethyl-4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline

20 7-Chloro-4-(3,5-dimethylpiperazin-1-yl)quinoline (100 mg, 0.36 mmol) and 4-fluorophenyl isocyanate (70 mg, 0.51 mmol) are reacted according to method C yielding the title product as colorless solid after column chromatography with CH_2Cl_2 -MeOH.

25 ^1H NMR ([D]₆-DMSO) δ 1.5 (d, 6H), 2.9 (m, 2H), 3.4 (m, 2H), 4.4 (m, 2H), 7.0 (m, 3H), 7.5 (m, 2H), 7.6 (dd, 1H), 8.0 (d, 1H), 8.2 (d, 1H), 8.4 (s, 1H), 8.7 (d, 1H).

Example 56

Preparation of 7-Chloro-4-[3,6-dimethyl-4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]quinoline, trifluoroacetate

30

Sodium hydride (72 mg, 3 mmol) is added to a solution of 7-chloro-4-quinolone (359 mg, 2 mmol) in DMF. After 1 h at 40 °C *N*-phenyl (trifluoromethylsulfon)imide (1.0 g,

2.8 mmol) and, 1 h later, 1-*tert*-butoxycarbonyl-2,5-dimethylpiperazine (1.8 g, 8 mmol) is added. The reaction mixture is stirred at 80 °C for 2 days, concentrated, diluted with EtOAc, washed with water and brine, dried (Na_2SO_4), and concentrated. The residue is purified by reversed phase HPLC with water-MeCN-TFA to give 4-[4-(*tert*-

5 butoxycarbonyl)-2,5-dimethylpiperazin-1-yl]-7-chloroquinoline (188 mg, 0.5 mmol) which is deprotected with TFA in CH_2Cl_2 and transformed after being washed with base into the product with 4-fluorophenyl isocyanate (31 μL , 0.27 mmol) according to method C giving the title product as a colorless solid.

10 ^1H NMR ($[\text{D}]_6\text{-DMSO}$) δ 1.1 (d, 3H), 1.3 (d, 3H), 3.6-3.8 (m, 3H), 4.1 (m, 1H), 4.5 (m, 2H), 7.0 (m, 2H), 7.3 (d, 1H), 7.4 (m, 2H), 7.7 (d, 1H), 8.0 (s, 1H), 8.2 (d, 1H), 8.6 (s, 1H), 8.7 (d, 1H).

Example 57

15 **Preparation of 7-Chloro-4-[5-(4-fluorophenylaminocarbonyl)-2,5-diazanorbornan-2-yl]quinoline, trifluoroacetate**

Preparation of 7-Chloro-4-(2,5-diazanorborn-2-yl)quinoline

20 4,7-Dichloroquinoline (0.38 g, 1.9 mmol) and 2,5-diazanorbornane dihydrobromide (1.0 g, 3.8 mmol), and DABCO (1.3 g, 11.4 mmol) in *n*-butanol (20 mL) are reacted according to method B yielding the product after column chromatography with CH_2Cl_2 -MeOH.

25 ^1H NMR (CDCl_3) δ 2.0 (dd, 2H), 3.2 (dd, 1H), 3.4 (d, 2H), 3.9 (s, 1H), 4.1 (dd, 2H), 4.5 (s, 1H), 6.4 (d, 1H), 7.3 (d, 1H), 7.9 (s, 1H), 7.95 (d, 1H), 8.5 (d, 1H).

25 *Preparation of 7-Chloro-4-[5-(4-fluorophenylaminocarbonyl)-2,5-diazanorbornan-2-yl]quinoline*

7-Chloro-4-(2,5-diazanorborn-2-yl)quinoline (110 mg, 0.42 mmol) and 4-fluorophenyl isocyanate (67 μL , 0.59 mmol) are reacted according to method C yielding the title product as a colorless solid after column chromatography with CH_2Cl_2 -MeOH.

30 ^1H NMR (CD_3OD) δ 2.2 (m, 2H), 3.3 (s, 1H), 3.7-4.1 (m, 4H), 5.0 (s, 1H), 7.0 (m, 3H), 7.3 (m, 3H), 7.6 (d, 1H), 7.9 (s, 1H), 8.3 (d, 1H), 8.4 (m, 1H).

Example 58**Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)-1,4-diaz pin-1-yl]quinoline**

5

Synthesis analogous to example 1.

¹H NMR (CDCl₃) δ 2.25 (m, 2H), 3.45 (m, 2H), 3.50 (m, 2H), 3.80 (t, 2H), 3.90 (m, 2H), 6.30 (s, 1H), 6.85 (d, 1H), 7.00 (t, 2H), 7.35 (m, 2H), 7.42 (d, 1H), 8.00 (d, 1H), 8.05 (s, 1H), 8.70 (d, 1H).

10

Example 59**Preparation of 7-Chloro-4-[4-(3,4-difluorophenylaminocarbonyl)piperazin-1-yl]quinoline**

15 7-Chloro-4-(piperazin-1-yl)quinoline (0.16 g, 0.65 mmol) and 3,4-difluorophenyl isocyanate (76 μL, 0.65 mmol) in THF (10 mL) are reacted according to method C yielding the title product as a colorless foam after column chromatography with hexane-EtOAc.

16 ¹H NMR ([D]₆-DMSO) δ 3.18 (m, 4H), 3.72 (m, 4H), 7.04 (d, 1H), 7.22 (m, 1H), 7.29 (q, 1H), 7.56 (dd, 1H), 7.62 (ddd, 2H), 7.98 (d, 1H), 8.09 (d, 1H), 8.70 (d, 1H), 8.85 (br. s, 1H).

Example 60**Preparation of 7-Chloro-4-[4-(2,3,4-trifluorophenylaminocarbonyl)piperazin-1-yl]quinoline**

25 7-Chloro-4-(piperazin-1-yl)quinoline (0.16 g, 0.65 mmol) and 2,3,4-difluorophenyl isocyanate (79 μL, 0.65 mmol) in THF (10 mL) are reacted according to method C yielding the title product as colorless solid after column chromatography with hexane-EtOAc.

30 ¹H NMR ([D]₆-DMSO) δ 3.18 (m, 4H), 3.72 (m, 4H), 7.04 (d, 1H), 7.22 (m, 2H), 7.56 (dd, 1H), 7.98 (d, 1H), 8.09 (d, 1H), 8.68 (br. s, 1H), 8.72 (d, 1H).

Example 61**Preparation of 7-Chloro-4-[4-(phenylaminocarbonyl)piperazin-1-yl]quinoline**

5 7-Chloro-4-(piperazin-1-yl)quinoline (62 mg, 0.25 mmol) and phenyl isocyanate (33 µL, 0.30 mmol) in THF (10 mL) are reacted according to method C yielding the title product as colorless needles after column chromatography with hexane-EtOAc.

10 ^1H NMR ($[\text{D}]_6\text{-DMSO}$) δ 3.18 (m, 4H), 3.72 (m, 4H), 6.92 (t, 1H), 7.04 (d, 1H), 7.22 (t, 2H), 7.46 (d, 2H), 7.57 (dd, 1H), 7.98 (d, 1H), 8.10 (d, 1H), 8.64 (br. s, 1H), 8.71 (d, 1H).

Example 62**Preparation of 7-Chloro-4-[4-(4-methylphenylaminocarbonyl)piperazin-1-yl]quinoline**

15 7-Chloro-4-(piperazin-1-yl)quinoline (62 mg, 0.25 mmol) and *p*-tolyl isocyanate (32 µL, 0.25 mmol) in CH_2Cl_2 (5 mL) are reacted according to method C yielding the title product as a colorless solid.

20 ^1H NMR ($[\text{D}]_6\text{-DMSO}$) δ 2.20 (s, 3H), 3.17 (m, 4H), 3.70 (m, 4H), 7.01 (d, 2H), 7.04 (d, 1H), 7.33 (d, 2H), 7.56 (dd, 1H), 7.98 (d, 1H), 8.08 (d, 1H), 8.53 (br. s, 1H), 8.70 (d, 1H).

Example 63**Preparation of 7-Chloro-4-[4-(3-methylphenylaminocarbonyl)piperazin-1-yl]quinoline**

25 7-Chloro-4-(piperazin-1-yl)quinoline (62 mg, 0.25 mmol) and *m*-tolyl isocyanate (32 µL, 0.25 mmol) in CH_2Cl_2 (5 mL) are reacted according to method C yielding the title product as colorless solid.

30 ^1H NMR ($[\text{D}]_6\text{-DMSO}$) δ 2.21 (s, 3H), 3.18 (m, 4H), 3.71 (m, 4H), 6.74 (d, 1H), 7.04 (d, 1H), 7.09 (t, 1H), 7.25 (d, 1H), 7.29 (d, 1H), 7.56 (dd, 1H), 7.98 (d, 1H), 8.09 (d, 1H), 8.56 (br. s, 1H), 8.71 (d, 1H).

Example 64**Preparation of 7-Chloro-4-[4-(2-methylphenylaminocarbonyl)piperazin-1-yl]quinoline**

5

7-Chloro-4-(piperazin-1-yl)quinoline (62 mg, 0.25 mmol) and *o*-tolyl isocyanate (31 μ L, 0.25 mmol) in CH₂Cl₂ (5 mL) are reacted according to method C yielding the title product as colorless solid.

10 ¹H NMR ([D]₆-DMSO) δ 2.17 (s, 3H), 3.20 (m, 4H), 3.70 (m, 4H), 7.02 (td, 1H), 7.05 (d, 1H), 7.11 (td, 1H), 7.16 (dd, 1H), 7.18 (dd, 1H), 7.29 (d, 1H), 7.56 (dd, 1H), 7.98 (d, 1H), 8.09 (d, 1H), 8.18 (br. s, 1H), 8.71 (d, 1H).

Example 65**Preparation of 4-[4-(4-*n*-Butylphenylaminocarbonyl)piperazin-1-yl]-7-chloroquinoline**

15 7-Chloro-4-(piperazin-1-yl)quinoline (62 mg, 0.25 mmol) and 4-(*n*-butyl)phenyl isocyanate (57 μ L, 0.30 mmol) in THF (10 mL) are reacted according to method C yielding the title product as a colorless solid after column chromatography with hexane-EtOAc.

20 ¹H NMR ([D]₆-DMSO) δ 0.88 (t, 3H), 1.25 (sext, 2H), 1.50 (pent, 2H), 2.50 (m, 2H), 3.17 (m, 4H), 3.71 (m, 4H), 7.03 (d, 2H), 7.04 (d, 1H), 7.34 (d, 2H), 7.56 (dd, 1H), 7.98 (d, 1H), 8.09 (d, 1H), 8.55 (br. s, 1H), 8.71 (d, 1H).

25 **Example 66**

Preparation of 7-Chloro-4-[4-(4-methoxyphenylaminocarbonyl)piperazin-1-yl]quinoline

30 7-Chloro-4-(piperazin-1-yl)quinoline (316 mg, 1.28 mmol) and 4-methoxyphenyl isocyanate (194 μ L, 1.50 mmol) in THF (10 mL) are reacted according to method C yielding the title product as a colorless needles after column chromatography with hexane-EtOAc and recrystallization from ether.

¹H NMR ([D]₆-DMSO) δ 3.18 (m, 4H), 3.68 (s, 3H), 3.71 (m, 4H), 7.01 (d, 2H), 7.04 (d, 1H), 7.34 (d, 2H), 7.56 (dd, 1H), 7.98 (d, 1H), 8.09 (d, 1H), 8.48 (br. s, 1H), 8.71 (d, 1H).

5 **Example 67**

Preparation of 4-[4-(4-Benzylxyloxyphenylaminocarbonyl)piperazin-1-yl]-7-chloroquinoline

7-Chloro-4-(piperazin-1-yl)quinoline (62 mg, 0.25 mmol) and 4-benzylxyloxyphenyl isocyanate (68 mg, 0.30 mmol) in THF (10 mL) are reacted according to method C yielding the title product as a yellow solid after column chromatography with hexane-EtOAc.

¹H NMR ([D]₆-DMSO) δ 3.18 (m, 4H), 3.71 (m, 4H), 5.02 (s, 2H), 6.89 (d, 2H), 7.04 (d, 1H), 7.28-7.42 (m, 7H), 7.56 (dd, 1H), 7.98 (d, 1H), 8.08 (d, 1H), 8.49 (br. s, 1H), 8.71 (d, 1H).

Example 68

Preparation of 4-[4-(Benzylaminocarbonyl)piperazin-1-yl]-7-chloroquinoline

7-Chloro-4-(piperazin-1-yl)quinoline (62 mg, 0.25 mmol) and benzyl isocyanate (37 μL, 0.30 mmol) in THF (10 mL) are reacted according to method C yielding the title product as a colorless oil after column chromatography with hexane-EtOAc.

¹H NMR ([D]₆-DMSO) δ 3.12 (m, 4H), 3.60 (m, 4H), 4.26 (d, 2H), 7.02 (d, 1H), 7.16-7.32 (m, 6H), 7.55 (dd, 1H), 7.96 (d, 1H), 8.07 (d, 1H), 8.70 (d, 1H).

25

Example 69

Preparation of 7-Chloro-4-[4-(phenethylaminocarbonyl)piperazin-1-yl]quinoline

7-Chloro-4-(piperazin-1-yl)quinoline (62 mg, 0.25 mmol) and phenethyl isocyanate (45 μL, 0.30 mmol) in THF (10 mL) are reacted according to method C yielding the title product as a colorless oil after column chromatography with hexane-EtOAc.

¹H NMR ([D]₆-DMSO) δ 2.71 (t, 2H), 3.10 (m, 4H), 3.26 (q, 2H), 3.57 (m, 4H), 6.74 (t,

1H), 7.01 (d, 1H), 7.10-7.30 (m, 5H), 7.55 (dd, 1H), 7.97 (d, 1H), 8.06 (d, 1H), 8.70 (d, 1H).

Example 70

5 **Preparation of 7-Chloro-4-[4-(phenylpropylaminocarbonyl)piperazin-1-yl]quinoline**

3-Phenyl-1-propylamine (71 μ L, 0.5 mmol), 4-nitrophenyl chloroformate (101 mg, 0.5 mmol), triethylamine (154 μ L, 1.1 mmol), and 7-chloro-4-(piperazin-1-yl)quinoline (124 mg, 0.5 mmol) are treated according to method D delivering the title product as a colorless oil after column chromatography with hexane-EtOAc.

^1H NMR ([D]₆-DMSO) δ 1.72 (pent, 2H), 2.58 (t, 2H), 3.05 (q, 2H), 3.11 (m, 4H), 3.55 (m, 4H), 6.63 (t, 1H), 7.01 (d, 1H), 7.14 (t, 1H), 7.19 (d, 2H), 7.26 (t, 2H), 7.54 (dd, 1H), 7.97 (d, 1H), 8.06 (d, 1H), 8.70 (d, 1H).

15

Example 71

Preparation of 7-Chloro-4-[4-[[methoxycarbonyl(phenyl)methyl]aminocarbonyl]piperazin-1-yl]quinoline

20 Phenylglycine methyl ester hydrochloride (606 mg, 3.0 mmol), 4-nitrophenyl chloroformate (606 mg, 3.0 mmol), triethylamine (1.84 mL, 13.2 mmol), and 7-chloro-4-(piperazin-1-yl)quinoline (743 mg, 3.0 mmol) are treated according to method D delivering the title product as a colorless solid after column chromatography with hexane-EtOAc.

25 ^1H NMR ([D]₆-DMSO) δ 3.12 (m, 4H), 3.60 (s, 3H), 3.64 (m, 4H), 5.32 (d, 1H), 7.02 (d, 1H), 7.26 (d, 1H), 7.31-7.42 (m, 5H), 7.54 (dd, 1H), 7.96 (d, 1H), 8.06 (d, 1H), 8.69 (d, 1H).

Example 72

30 **Preparation of 7-Chloro-4-[4-(1-phenylethylaminocarbonyl)piperazin-1-yl]quinoline**

7-Chloro-4-(piperazin-1-yl)quinoline (62 mg, 0.25 mmol) and α -methylbenzyl isocyanate (42 μ L, 0.30 mmol) in THF (10 mL) are reacted according to method C yielding the title product as a colorless solid after column chromatography with hexane-EtOAc.

5 ^1H NMR ($[\text{D}]_6\text{-DMSO}$) δ 1.36 (d, 3H), 3.12 (m, 4H), 3.59 (m, 4H), 4.84 (pent, 1H), 6.92 (br. d, 1H), 7.02 (d, 1H), 7.16 (t, 1H), 7.28 (m, 4H), 7.54 (dd, 1H), 7.96 (d, 1H), 8.06 (d, 1H), 8.70 (d, 1H).

Example 73

10 **Preparation of 7-Chloro-4-[4-(2-chlorobenzylaminocarbonyl)piperazin-1-yl]quinoline**

2-Chlorobenzylamine (60 μ L, 0.5 mmol), 4-nitrophenyl chloroformate (101 mg, 0.5 mmol), triethylamine (238 μ L, 1.7 mmol), and 7-chloro-4-(piperazin-1-yl)quinoline (124 mg, 0.5 mmol) are treated according to method D delivering 147 mg of the product as a pale yellow solid after column chromatography with hexane-EtOAc.

^1H NMR (CDCl_3) δ 3.19 (m, 4H), 3.65 (m, 4H), 4.55 (d, 2H), 5.07 (t, 1H), 6.82 (d, 1H), 7.24 (m, 2H), 7.36 (m, 1H), 7.44 (m, 2H), 7.92 (d, 1H), 8.04 (d, 1H), 8.72 (d, 1H).

20 **Example 74**

Preparation of 7-Chloro-4-[4-(2-methylbenzylaminocarbonyl)piperazin-1-yl]quinoline

2-Methylbenzylamine (62 μ L, 0.5 mmol), 4-nitrophenyl chloroformate (101 mg, 0.5 mmol), triethylamine (238 μ L, 1.7 mmol), and 7-chloro-4-(piperazin-1-yl)quinoline (124 mg, 0.5 mmol) are treated according to method D delivering the title product as a pale yellow solid after column chromatography with hexane-EtOAc.

^1H NMR (CDCl_3), δ 3.20 (m, 4H), 3.66 (m, 4H), 4.46 (d, 2H), 4.60 (t, 1H), 6.83 (d, 1H), 7.19 (m, 3H), 7.28 (m, 1H), 7.44 (dd, 1H), 7.94 (d, 1H), 8.05 (d, 1H), 8.73 (d, 1H).

30

Example 75

Preparation of 7-Chloro-4-[4-(2-trifluoromethylbenzylaminocarbonyl)piperazin-1-

yl]quinolin

2-Trifluoromethylbenzylamine (70 μ L, 0.5 mmol), 4-nitrophenyl chloroformate (101 mg, 0.5 mmol), triethylamine (238 μ L, 1.7 mmol), and 7-chloro-4-(piperazin-1-yl)quinoline (124 mg, 0.5 mmol) are treated according to method D delivering the title product as a pale yellow solid after column chromatography with hexane-EtOAc.

⁵ 1 H NMR ($CDCl_3$) δ 3.20 (m, 4H), 3.66 (m, 4H), 4.64 (d, 2H), 4.86 (t, 1H), 6.92 (d, 1H), 7.38 (t, 1H), 7.43 (dd, 1H), 7.54 (t, 1H), 7.62 (d, 1H), 7.64 (d, 1H), 7.92 (d, 1H), 8.04 (d, 1H), 8.72 (d, 1H).

10

Example 76**Preparation of 7-Chloro-4-[4-(1-indanylaminocarbonyl)piperazin-1-yl]quinoline**

2-Aminoindane (32 μ L, 0.25 mmol), 4-nitrophenyl chloroformate (51 mg, 0.25 mmol), ¹⁵ triethylamine (140 μ L, 1.0 mmol), and 7-chloro-4-(piperazin-1-yl)quinoline (62 mg, 0.25 mmol) are treated according to method D delivering the title product as a colorless solid after column chromatography with hexane-EtOAc.

¹⁶ 1 H NMR ($CDCl_3$) δ 1.85 (m, 1H), 2.67 (m, 1H), 2.90 (m, 1H), 3.00 (m, 1H), 3.22 (m, 4H), 3.67 (m, 4H), 4.72 (d, 1H), 5.45 (q, 1H), 6.84 (d, 1H), 7.23 (m, 3H), 7.34 (d, 1H), ²⁰ 7.44 (dd, 1H), 7.94 (d, 1H), 8.05 (d, 1H), 8.73 (d, 1H).

25

Example 77**Preparation of 7-Chloro-4-[4-(1,2,3,4-tetrahydronaphth-1-ylaminocarbonyl)piperazin-1-yl]quinoline**

25

1,2,3,4-Tetrahydronaphthylamine (36 μ L, 0.25 mmol), 4-nitrophenyl chloroformate (51 mg, 0.25 mmol), triethylamine (140 μ L, 1.0 mmol), and 7-chloro-4-(piperazin-1-yl)quinoline (62 mg, 0.25 mmol) are treated according to method D delivering the title product as a colorless oil after column chromatography with hexane-EtOAc.

³⁰ 1 H NMR ($CDCl_3$) δ 1.88 (m, 3H), 2.08 (m, 1H), 2.80 (m, 2H), 3.20 (m, 4H), 3.66 (m, 4H), 4.72 (d, 1H), 5.12 (m, 1H), 6.84 (d, 1H), 7.10 (dd, 1H), 7.18 (d, 1H), 7.20 (d, 1H), 7.34 (dd, 1H), 7.44 (dd, 1H), 7.92 (d, 1H), 8.04 (d, 1H), 8.72 (d, 1H).

Example 78**Preparation of 7-Chloro-4-[4-(4-pyridinylaminocarbonyl)piperazin-1-yl]quinoline**

5 At 0 °C under nitrogen, to a solution of 4-aminopyridine (114 mg, 1.21 mmol) and 4-nitrophenyl chloroformate (244 mg, 1.21 mmol) in dichloromethane (5 mL) is added diisopropyl(ethyl)amine dropwise (391 mg, 3.03 mmol). After 10 min, the ice bath is removed, and the reaction mixture is stirred at rt. for an additional hour. After removal of solvent under vacuum, EtOAc (5 mL), diisopropyl(ethyl)amine (261 mg, 2.02 mmol)

10 and 7-chloro-4-(piperazin-1-yl)quinoline (250 mg, 1.01 mmol) are added. The resulting mixture is kept at reflux for about 5 hr, then cooled to rt, diluted with EtOAc (10 mL), and quenched with saturated sodium bicarbonate (5 mL). The aqueous solution is extracted with EtOAc (2x10 mL), and the combined organic phase is washed with brine (10 mL) and dried over sodium sulfate. Removal of solvent followed by

15 purification through flash chromatography with CH₂Cl₂-MeOH affords the title product.
¹H NMR (CDCl₃) δ 8.61 (d, 2H), 8.32 (d, 2H), 7.98 (s, 1H), 7.86 (d, 1H), 7.8 (d, 2H), 7.74 (d, 1H), 3.79 (br. s, 4H), 3.12 (br. s, 4H).

Example 79

20 **Preparation of 7-Chloro-4-[4-(3-pyridinylaminocarbonyl)piperazin-1-yl]quinoline**

As described for example 78, 3-aminopyridine (114 mg, 1.21 mmol), 4-nitrophenyl chloroformate (244 mg, 1.21 mmol), diisopropyl(ethyl)amine (650 mg, 5.05 mmol), and 7-chloro-4-(piperazin-1-yl)quinoline (250 mg, 1.01 mmol) are reacted affording the title

25 product after flash chromatography with CH₂Cl₂-MeOH.

¹H NMR (DSMO-d₆/TFA) δ 9.64 (s, 1H), 9.13 (s, 1H), 8.69 (d, 1H), 8.46 -8.53 (m, 2H), 8.25 (d, 1H), 8.07 (d, 1H), 7.95 (dd, 1H), 7.68 (dd, 1H), 7.19 (d, 1H), 4.02 (m, 4H), 3.81 (m, 4H).

30 **Example 80**

Preparation of 7-Chloro-4-[4-(2-pyridinylaminocarbonyl)piperazin-1-yl]quinoline

As described for example 78, 2-aminopyridine (142 mg, 1.51 mmol), 4-nitrophenyl chloroformate (304 mg, 1.51 mmol), diisopropyl(ethyl)amine (650 mg, 5.05 mmol), and 7-chloro-4-(piperazin-1-yl)quinoline (250 mg, 1.01 mmol) are reacted affording the title product after flash chromatography with CH₂Cl₂-MeOH.

5 ¹H NMR (DSMO-d₆/TFA) δ 8.62 (d, 1H), 8.18 (m, 2H), 7.96 (s, 1H), 7.83 (d, 1H), 7.38 (d, 1H), 7.22 (d, 1H), 6.78 (d, 1H), 3.91 (m, 2H), 3.82 (m, 2H), 3.2 (m, 4H).

Example 81

Preparation of 7-Chloro-4-[4-(4-trifluoromethylphenylaminocarbonyl)piperazin-1-yl]quinoline

At room temp. under nitrogen, to a solution of 7-chloro-4-(piperazin-1-yl)quinoline (200 mg, 0.81 mmol) and diisopropyl(ethyl)amine (209 mg, 1.62 mmol) in acetonitrile (5 mL) is added 4-trifluoromethylphenyl isocyanate (197 mg, 0.97 mmol). The reaction mixture is stirred at room temp. for 2 h, concentrated, and the residue is purified by flash chromatography with CH₂Cl₂-MeOH to afford the title product.

15 ¹H NMR (DSMO-d₆/TFA) δ 9.65 (s, 1H), 8.83 (d, 1H), 8.32 (d, 1H), 8.01 (d, 1H), 7.53-7.74 (m, 4H), 7.07 (d, 1H), 4.16 (m, 8H).

20 **Example 82**

Preparation of 7-Chloro-4-[4-(4-ethoxyphenylaminocarbonyl)piperazin-1-yl]quinoline

As described for 7-chloro-4-[4-(4-trifluoromethylphenylaminocarbonyl)piperazin-1-yl]quinoline, 7-chloro-4-(piperazin-1-yl)quinoline (150 mg, 0.61 mmol), diisopropyl(ethyl)amine (158 mg, 1.22 mmol), and 4-ethoxyphenyl isocyanate (119 mg, 0.73 mmol) are reacted to give the title product after flash chromatography with CH₂Cl₂-MeOH.

25 ¹H NMR (DSMO-d₆/TFA) δ 8.65 (d, 1H), 8.46 (s, 1H), 8.42 (d, 1H), 7.98 (d, 1H), 7.67 (dd, 1H), 7.26-7.35 (m, 3H), 7.16 (d, 1H), 6.81 (m, 3H), 3.97 (m, 6H), 3.76 (m, 4H), 1.24 (t, 3H);

Example 83**Preparation of 7-Chloro-4-[4-(4-isopropylphenylaminocarbonyl)piperazin-1-yl]quinoline**

5 As described for 7-chloro-4-[4-(4-trifluoromethylphenylaminocarbonyl)piperazin-1-yl]quinoline, 7-chloro-4-(piperazin-1-yl)quinoline (150 mg, 0.61 mmol), diisopropyl(ethyl)amine (158 mg, 1.22 mmol), and 4-isopropylphenyl isocyanate (117 mg, 0.73 mmol) are reacted to give the title product after flash chromatography with CH₂Cl₂-MeOH.

10 ¹H NMR (DSMO-d₆/TFA) δ 8.65 (d, 1H), 8.58 (s, 1H), 8.23 (d, 1H), 7.66 (d, 1H), 7.37 (d, 2H), 7.18 (d, 1H), 7.06 (d, 2H), 3.92 (m, 4H), 3.76 (m, 4H), 2.77 (m, 1H), 1.18 (d, 6H).

Example 84

15 **Preparation of 4-[4-(4-Acetylphenylaminocarbonyl)piperazin-1-yl]-7-chloroquinoline**

As described for 7-chloro-4-[4-(4-trifluoromethylphenylaminocarbonyl)piperazin-1-yl]quinoline, 7-chloro-4-(piperazin-1-yl)quinoline (250 mg, 1.01 mmol), diisopropyl(ethyl)amine (261 mg, 2.02 mmol), and 4-acetylphenyl isocyanate (195 mg, 1.21 mmol) are reacted to give the title product after flash chromatography with CH₂Cl₂-MeOH.

20 ¹H NMR (DSMO-d₆/TFA), δ 9.07 (s, 1H), 8.69 (d, 1H), 8.23 (d, 1H), 8.02 (s, 1H), 7.83 (d, 2H), 7.65 (d, 1H), 7.61 (m, 2H), 7.19 (d, 1H), 3.99 (m, 4H), 3.78 (m, 4H), 2.5 (s, 25 3H).

Example 85**Preparation of 7-Chloro-4-[4-(4-dimethylaminophenylaminocarbonyl)piperazin-1-yl]quinoline**

30 As described for 7-chloro-4-[4-(4-trifluoromethylphenylaminocarbonyl)piperazin-1-yl]quinoline, 7-chloro-4-(piperazin-1-yl)quinoline (200 mg, 0.81 mmol), diisopropyl(ethyl)amine (209 mg, 1.62 mmol), and 4-(dimethylamino)phenyl isocyanate

(157 mg, 0.97 mmol) are reacted to give the title product after flash chromatography with CH₂Cl₂-MeOH.

¹H NMR (DSMO-d₆/TFA) δ 9.64 (s, 1H), 9.12 (s, 1H), 8.68 (d, 1H), 8.52 (m, 2H), 8.24 (d, 1H), 8.03 (d, 1H), 7.94 (dd, 1H), 7.69 (dd, 1H), 7.18 (d, 1H), 4.02 (m, 4H), 3.82 (m, 5 H) 2.42 (s, 6H).

Example 86

Preparation of 7-Chloro-4-[4-(cyclopentylaminocarbonyl)piperazin-1-yl]quinoline

As described for 7-chloro-4-[4-(4-trifluoromethylphenylaminocarbonyl)piperazin-1-yl]quinoline, 7-chloro-4-(piperazin-1-yl)quinoline (180 mg, 0.73 mmol), diisopropyl(ethyl)amine (189 mg, 1.46 mmol), and cyclopentyl isocyanate (97 mg, 0.88 mmol) are reacted to give the title product after flash chromatography with CH₂Cl₂-MeOH.

¹H NMR (DSMO-d₆/TFA) δ 8.62 (d, 1H), 8.21 (d, 1H), 7.98 (s, 1H), 7.64 (d, 1H), 7.17 (d, 1H), 3.86 (m, 5H), 3.58 (m, 4H), 1.78 (m, 2H), 1.62 (m, 2H), 1.32-1.56 (m, 4H).

Example 87

Preparation of 7-Chloro-4-[4-(3-piperonylaminocarbonyl)piperazin-1-yl]quinoline

As described for 7-chloro-4-[4-(4-trifluoromethylphenylaminocarbonyl)piperazin-1-yl]quinoline, 7-chloro-4-(piperazin-1-yl)quinoline (180 mg, 0.73 mmol), diisopropyl(ethyl)amine (189 mg, 1.46 mmol), and 3,4-(methylenedioxy)phenyl isocyanate (143 mg, 0.88 mmol) are reacted to give the product after flash chromatography with CH₂Cl₂-MeOH.

¹H NMR (DSMO-d₆/TFA) δ 8.64 (d, 1H), 8.42 (s, 1H), 8.21 (d, 1H), 7.95 (d, 1H), 7.66 (dd, 1H), 7.15 (dd, 1H), 7.07 (d, 1H), 6.74(d, 1H), 5.83 (S, 2H), 4.41 (m, 4H), 4.2 (m, 4H).

Example 88

Preparation of 7-Chloro-4-[4-[4-(6-methylbenzothiazol-2-yl)phenylaminocarbonyl]piperazin-1-yl]quinoline

As described for 7-chloro-4-[4-(4-trifluoromethylphenylaminocarbonyl)piperazin-1-yl]quinoline, 7-chloro-4-(piperazin-1-yl)quinoline (120 mg, 0.49 mmol), diisopropyl(ethyl)amine (127 mg, 0.98 mmol), and 4-(6-methyl-2-benzothiazolyl)phenyl 5 isocyanate (100 mg, 0.38 mmol) are reacted to give the title product after flash chromatography with CH₂Cl₂-MeOH.

¹H NMR (DSMO-d₆/TFA) δ 8.97 (s, 1H), 8.62 (d, 1H), 8.23 (d, 1H), 7.96 (d, 1H), 7.92 (d, 2H), 7.8 (m, 2H), 7.64-7.71 (m, 3H), 7.25 (dd, 1H), 7.16 (d, 1H), 3.96 (m, 4H), 3.79 (m, 4H), 2.38 (s, 3H).

10

Example 89

Preparation of 7-Chloro-4-[4-(cyclobutylaminocarbonyl)piperazin-1-yl]quinoline

As described for example 78, cyclobutylamine (87 mg, 1.21 mmol), 4-nitrophenyl 15 chloroformate (244 mg, 1.21 mmol), diisopropyl(ethyl)amine (2 x 261 mg, 2 x 2.02 mmol), and 7-chloro-4-(piperazin-1-yl)quinoline (250 mg, 1.01 mmol) are reacted affording the title product after flash chromatography with CH₂Cl₂-MeOH.

¹H NMR (DSMO-d₆/TFA) δ 8.65 (d, 1H), 8.04 (d, 1H), 7.94 (s, 1H), 7.57 (d, 1H), 7.0 (d, 1H), 6.77 (d, 1H), 4.12 (m, 1H), 3.58 (m, 4H), 3.09 (m, 4H), 2.11 (m, 2H), 1.95 (m, 2H), 20 1.52 (m, 2H).

Example 90

Preparation of 7-Chloro-4-[4-(cyclohexylaminocarbonyl)piperazin-1-yl]quinoline

25 As described for 7-chloro-4-[4-(4-trifluoromethylphenylaminocarbonyl)piperazin-1-yl]quinoline, 7-chloro-4-(piperazin-1-yl)quinoline (200 mg, 0.81 mmol), diisopropyl(ethyl)amine (125 mg, 0.97 mmol), and cyclohexyl isocyanate (121 mg, 0.97 mmol) are reacted to give the title product after flash chromatography with CH₂Cl₂-MeOH.

30 ¹H NMR (DSMO-d₆/TFA) δ 8.62 (d, 1H), 8.21 (d, 1H), 7.96 (s, 1H), 7.63 (d, 1H), 7.16 (d, 1H), 3.92 (m, 4H), 3.61 (m, 4H), 3.41 (m, 1H), 1.4-1.8 (m, 4H), 1.0-1.3 (m, 6H).

Example 91**Preparation of 4-[4-(Adamant-1-ylaminocarbonyl)piperazin-1-yl]-7-chloroquinoline**

5 7-Chloro-4-(piperazin-1-yl)quinoline (124 mg, 0.5 mmol) and 1-adamantyl isocyanate (89 mg, 0.5 mmol) in CH₂Cl₂ (5 mL) are reacted according to method C yielding the title product as a colorless solid after column chromatography with hexane-EtOAc.

10 ¹H NMR ([D]₆-DMSO) δ 1.59 (m, 6H), 1.91 (m, 6H), 1.97 (m, 3H), 3.08 (m, 4H), 3.50 (m, 4H), 5.78 (br. s, 1H), 7.00 (d, 1H), 7.53 (dd, 1H), 7.96 (d, 1H), 8.04 (d, 1H), 8.68 (d, 1H).

Example 92**Preparation of 7-Chloro-4-[4-(exo-norborn-2-ylaminocarbonyl)piperazin-1-yl]quinoline**

15 exo-2-Aminonorbornane (59 µL, 0.5 mmol), 4-nitrophenyl chloroformate (101 mg, 0.5 mmol), triethylamine (140 µL, 1.0 mmol), and 7-chloro-4-(piperazin-1-yl)quinoline (124 mg, 0.5 mmol) are treated according to method D delivering the title product as a colorless foam after column chromatography with hexane-EtOAc.

20 ¹H NMR ([D]₆-DMSO) δ 1.05 (m, 3H), 1.37 (m, 3H), 1.50 (m, 2H), 2.07 (m, 1H), 2.15 (m, 1H), 3.10 (m, 4H), 3.42 (m, 1H), 3.57 (m, 4H), 6.33 (d, 1H), 7.00 (d, 1H), 7.54 (dd, 1H), 7.96 (d, 1H), 8.05 (d, 1H), 8.68 (d, 1H).

Example 93**Preparation of 7-Chloro-4-[4-(4-fluorophenylsulfonyl)piperazin-1-yl]quinoline**

7-Chloro-4-(piperazin-1-yl)quinoline (0.25 g, 1.0 mmol), 4-fluorobenzenesulfonyl chloride (0.23 g, 1.2 mmol), and triethylamine (349 µL, 2.5 mmol) are stirred at room temp. for 15 h. The reaction mixture is diluted with EtOAc, washed with water, dried (MgSO₄), and concentrated. The residue is purified by column chromatography with hexane –EtOAc yielding the title product as a colorless solid.

¹H NMR (CDCl₃) δ 3.31 (m, 8H), 6.84 (d, 1H), 7.28 (dd, 2H), 7.37 (dd, 1H), 7.75 (d,

1H), 7.85 (dd, 2H), 8.03 (d, 1H), 8.73 (d, 1H).

Example 94

7-Chloro-4-[4-(4-fluorophenylaminothiocarbonyl)piperazin-1-yl]quinoline

5

7-Chloro-4-(piperazin-1-yl)quinoline (0.20 g, 0.8 mmol) and 4-fluorophenyl isothiocyanate (0.14 g, 0.9 mmol) in THF (10 mL) are reacted according to method C yielding the title product as a solid after column chromatography with hexane-EtOAc.

10 ^1H NMR ($[\text{D}]_6\text{-DMSO}$) δ 3.29 (m, 4H), 4.18 (m, 4H), 7.04 (d, 1H), 7.12 (t, 2H), 7.30 (m, 2H), 7.56 (dd, 1H), 7.98 (d, 1H), 8.13 (d, 1H), 8.72 (d, 1H), 9.44 (s, 1H).

Example 95

7-Chloro-4-[4-[cyanimino(phenoxy)methyl]piperazin-1-yl]quinoline

15 7-Chloro-4-(piperazin-1-yl)quinoline (798 mg, 3.22 mmol) and diphenyl cyanocarbonimidate (853 mg, 3.58 mmol) in THF (10 mL) are stirred at room temp. for 15 h. The reaction mixture is diluted with ether, and the precipitate is filtered off to yield the title product as a colorless solid.

16 ^1H NMR (CDCl_3), δ 3.29 (m, 4H), 4.02 (m, 4H), 6.87 (d, 1H), 7.12 (d, 2H), 7.28 (t, 1H),
20 7.42 (t, 2H), 7.46 (dd, 1H), 7.91 (d, 1H), 8.06 (d, 1H), 8.76 (d, 1H).

Example 96**Preparation of 7-Chloro-4-[4-[cyanimino(ethoxy)methyl]piperazin-1-yl]quinoline**

7-Chloro-4-[4-(cyanimino(phenoxy)methyl)piperazin-1-yl]quinoline (196 mg, 0.5 mmol)

5 and potassium *tert*-butoxide (62 mg, 0.55 mmol) in EtOH (5 mL) are stirred at room temp. for 4 h. The reaction mixture is concentrated, and the residue purified by column chromatography delivering the title product.

¹H NMR (CDCl₃) δ 1.38 (t, 3H), 3.23 (m, 4H), 3.98 (m, 4H), 4.50 (q, 2H), 6.83 (d, 1H), 7.45 (dd, 2H), 7.89 (d, 1H), 8.05 (d, 1H), 8.74 (d, 1H).

10

Example 97**Preparation of 7-Chloro-4-[4-(cyanimino(4-fluorophenylamino)methyl)piperazin-1-yl]quinoline**

15 Diphenyl cyanocarbonimidate (0.24 g, 1.0 mmol) and 4-fluoroaniline (104 μL, 1.1 mmol) are stirred in isopropanol (10 mL) for 15 h. The reaction mixture is concentrated, and the residue dissolved in EtOAc. Washing with sat NaHCO₃, drying (MgSO₄) and removing the solvent delivers 0.34 g of N-cyano-N'-(4-fluorophenyl)-O-phenylisourea as a colorless solid. It is dissolved in pyridine (15 mL) and 7-chloro-4-(piperazin-1-yl)quinoline (0.25 g, 1.0 mmol) is added. The reaction mixture is heated at reflux for 4 h and afterwards concentrated. Column chromatography of the residue with hexane-EtOAc gives the title product as a colorless solid.

20 ¹H NMR ([D]₆-DMSO) δ 3.23 (m, 4H), 3.76 (m, 4H), 7.02 (d, 1H), 7.16 (m, 4H), 7.56 (dd, 1H), 7.98 (d, 1H), 8.07 (d, 1H), 8.72 (d, 1H), 9.45 (br. s, 1H).

25

Example 98**Preparation of 7-Chloro-4-[4-(cyanimino(4-fluorobenzylamino)methyl)piperazin-1-yl]quinoline**

30 7-Chloro-4-[4-(cyanimino(phenoxy)methyl)piperazin-1-yl]quinoline (107 mg, 0.27 mmol) and 4-fluorobenzylamine (62 μL, 0.54 mmol) are heated in pyridine (10 mL) for 3 h at reflux. The solvent is removed, and the residue purified by column

chromatography with CH₂Cl₂-MeOH giving the title product as a colorless solid.

¹H NMR (CDCl₃) δ 3.25 (m, 4H), 3.77 (m, 4H), 4.57 (d, 2H), 5.31 (t, 1H), 6.83 (d, 1H), 7.04 (t, 2H), 7.30 (dd, 2H), 7.44 (dd, 1H), 7.88 (d, 1H), 8.05 (d, 1H), 8.73 (d, 1H).

5 **Example 99**

Preparation of 4-[4-(*tert*-Butylamino(cyanimino)methyl)piperazin-1-yl]-7-chloroquinoline

7-Chloro-4-[4-(cyanimino(phenoxy)methyl)piperazin-1-yl]quinoline (53 mg, 0.14 mmol)

10 and *tert*-butylamine (2.0 mL, 19 mmol) are heated in pyridine (10 mL) for 6 h at reflux.

The solvent is removed, and the residue purified by column chromatography with CH₂Cl₂-MeOH giving the title product as a colorless solid.

¹H NMR (CDCl₃) δ 1.25 (s, 9H), 2.78 (m, 4H), 4.02 (m, 4H), 6.23 (d, 1H), 7.27 (dd, 2H), 7.80 (d, 1H), 8.53 (d, 1H), 8.69 (d, 1H).

15

Example 100

Preparation of 7-Chloro-4-[4-[cyanimino(ethylamino)methyl]piperazin-1-yl]quinoline

20 7-Chloro-4-[4-(cyanimino(phenoxy)methyl)piperazin-1-yl]quinoline (0.39 g, 1.0 mmol), ethylamine hydrochloride (1.68 g, 20.6 mmol), and diaza[2.2.2]bicyclooctane (2.24 g, 20.0 mmol) in isopropanol (20 mL) are heated in a sealed tube for 4 h at 110 °C. The reaction mixture is concentrated, and the residue is suspended in EtOAc. Washing with water, drying (MgSO₄), and removing the solvent leads to a colorless solid, which 25 is washed with hot ether to yield the title product.

¹H NMR ([D]₆-DMSO) δ 1.13 (t, 3H), 3.20 (m, 4H), 3.35 (m, 2H), 3.68 (m, 4H), 7.01 (d, 1H), 7.39 (br. t, 1H), 7.55 (dd, 1H), 7.97 (d, 1H), 8.06 (d, 1H), 8.70 (d, 1H).

Example 101

30 **Preparation of 7-Chloro-4-[4-[4-(4-fluorophenylamino]-2,3-dioxocyclobut-1-enyl]piperazin-1-yl]quinoline**

4-(4-Fluorophenylamino)-1-methoxy-2,3-dioxocyclobut-1-ene

Dimethyl squarate (142 mg, 1.0 mmol) and 4-fluoroaniline (104 μ L, 1.1 mmol) in isopropanol (10 mL) are stirred for 60 h at room temp. The reaction mixture is diluted with EtOAc, washed with water, dried ($MgSO_4$), and concentrated giving the desired product.

1H NMR ([D]₆-DMSO) δ 4.36 (s, 3H), 7.18 (t, 2H), 7.28 (br. m, 1H).

7-Chloro-4-[4-(4-fluorophenylamino)-2,3-dioxocyclobut-1-enyl]piperazin-1-yl]quinoline

7-Chloro-4-(piperazin-1-yl)quinoline (247 mg, 1.0 mmol) and 4-(4-fluorophenylamino)-1-methoxy-2,3-dioxocyclobut-1-ene (218 mg, 1.0 mmol) in pyridine (20 mL) are heated at reflux for 4 h. The reaction mixture is concentrated, and the residue purified by column chromatography with CH_2Cl_2 -MeOH and recrystallization from ether yielding the title product as a yellow solid.

1H NMR ([D]₆-DMSO) δ 3.31 (m, 4H), 4.03 (m, 4H), 7.06 (d, 1H), 7.15 (t, 2H), 7.24 (dd, 2H), 7.57 (dd, 1H), 7.99 (d, 1H), 8.10 (d, 1H), 8.73 (d, 1H).

Example 102**Preparation of 4-[4-(Benzimidazol-2-yl)piperazin-1-yl]-7-chloroquinoline**

7-Chloro-4-(piperazin-1-yl)quinoline (247 mg, 1.0 mmol) and 2-chlorobenzimidazole (46 mg, 0.3 mmol) are heated at 150 °C for 1 h. The mixture is dissolved in EtOAc-water. The organic layer is separated, dried ($MgSO_4$), and concentrated. Column chromatography of the residue with hexane-EtOAc furnishes the title product as colorless solid.

1H NMR ([D]₆-DMSO) δ 3.30 (m, 4H), 3.78 (m, 4H), 6.88 (t, 1H), 6.94 (t, 1H), 7.07 (d, 1H), 7.17 (d, 1H), 7.22 (d, 1H), 7.58 (dd, 1H), 7.98 (d, 1H), 8.12 (d, 1H), 8.71 (d, 1H).

Example 103**Preparation of 7-Chloro-4-[4-(pyrimidin-2-yl)piperazin-1-yl]-7-chloroquinoline**

4,7-Dichloroquinoline (0.10 g, 0.5 mmol) and 1-(pyrimidin-2-yl)piperazine (0.33 g, 2.0 mmol) are heated at 150 °C for 2.5 h. The mixture is dissolved in EtOAc-water. The organic layer is separated, dried (MgSO_4), and concentrated. Column chromatography of the residue with hexane-EtOAc furnishes the title product as a colorless solid.

5 ^1H NMR ($[\text{D}]_6\text{-DMSO}$) δ 3.23 (m, 4H), 4.00 (m, 4H), 6.66 (t, 1H), 7.03 (d, 1H), 7.57 (dd, 1H), 7.98 (d, 1H), 8.12 (d, 1H), 8.38 (d, 2H), 8.70 (d, 1H).

Example 104

Preparation of 7-Chloro-4-[4-[ethoxyimino(4-fluorophenylamino)methyl]piperazin-1-yl]quinoline

Preparation of N-Ethoxy-N'-(4-fluorophenyl)thiourea

4-Fluorophenyl isothiocyanate (306 mg, 2.0 mmol), O-ethyl hydroxylamine hydrochloride (585 mg, 6.0 mmol), and triethylamine (1.5 mL, 11.0 mmol) are stirred in 15 CH_2Cl_2 (50 mL) for 2 h. The reaction mixture is diluted with ether, washed with water and 1 M HCl, dried (MgSO_4), and concentrated. The residue is purified by column chromatography with hexane –EtOAc yielding a colorless solid.

1 ^1H NMR (CDCl_3) δ 1.15 (t, 3H), 4.05 (q, 2H), 7.06 (dd, 2H), 7.51 (dd, 2H), 8.41 (br. s, 1H).

20

Preparation of 7-Chloro-4-[4-(ethoxyimino(4-fluorophenylamino)methyl)piperazin-1-yl]quinoline

7-Chloro-4-(piperazin-1-yl)quinoline (124 mg, 0.5 mmol), *N*-ethoxy-*N'*-(4-fluorophenyl)thiourea (105 mg, 0.49 mmol), and sodium periodate (118 mg, 0.55 mmol) in DMF-water 1:1 (20 mL) are heated at 85 °C for 1 h. The reaction mixture is diluted with water and extracted with ether. The organic phase is dried (MgSO_4) and concentrated. The residue is purified by column chromatography with hexane –EtOAc yielding the title product as pale yellow crystals.

30 ^1H NMR (CDCl_3) δ 1.29 (t, 3H), 3.17 (m, 4H), 3.31 (m, 4H), 4.03 (q, 2H), 6.58 (br. s, 1H), 6.79 (d, 1H), 6.96 (t, 2H), 7.15 (dd, 2H), 7.39 (dd, 1H), 7.89 (d, 1H), 8.01 (d, 1H), 8.69 (d, 1H).

Example 105**Preparation of 7-Chloro-4-[4-[methoxyimino(4-fluorophenylamino)methyl]piperazin-1-yl]quinoline**

5

Preparation of N-(4-Fluorophenyl)-N'-methoxythiourea

4-Fluorophenyl isothiocyanate (306 mg, 2.0 mmol), O-methyl hydroxylamine hydrochloride (835 mg, 10.0 mmol), and triethylamine (1.5 mL, 11.0 mmol) are stirred in THF-water (6 mL) for 2 h. The reaction mixture is diluted with ether, washed with

10 water, dried (MgSO_4), and concentrated yielding 240 mg of pale yellow crystals.

^1H NMR (CDCl_3) δ 3.85 (s, 3H), 7.06 (dd, 2H), 7.52 (dd, 2H, 8.41 (br. s, 1H).

7-Chloro-4-[4-(methoxyimino(4-fluorophenylamino)methyl)piperazin-1-yl]quinoline

7-Chloro-4-(piperazin-1-yl)quinoline (297 mg, 1.2 mmol), *N*-(4-Fluorophenyl)-*N'*-

methoxythiourea (240 mg, 1.2 mmol), and sodium periodate (278 mg, 1.3 mmol) in

15 DMF-water 1:1 (40 mL) are heated at 85 °C for 1 h. The reaction mixture is diluted with water and extracted with ether. The organic phase is dried (MgSO_4) and concentrated. The residue is purified by column chromatography with hexane –EtOAc yielding the title product.

20 ^1H NMR (CDCl_3) δ 3.14 (m, 4H), 3.30 (m, 4H), 3.79 (s, 3H), 6.54 (br. s, 1H), 6.80 (d, 1H), 6.96 (t, 2H), 7.14 (dd, 2H), 7.38 (dd, 1H), 7.88 (d, 1H), 8.01 (d, 1H), 8.69 (d, 1H).

Example 106**Preparation of 7-Chloro-4-[4-(4-fluorobenzylamino(imino)methyl)piperazin-1-yl]quinoline***7-Chloro-4-(4-cyanopiperazin-1-yl)quinoline*

A suspension of 7-Chloro-4-(piperazin-1-yl)quinoline (496 mg, 2.0 mmol) and NaHCO_3 (504 mg, 6.0 mmol) in EtOH is treated with cyanogen bromide (5 M in MeCN, 440 μL ,

30 2.2 mmol) and stirred at room temp. for 2 h. The solvent is removed, and the residue suspended in CH_2Cl_2 and filtered. The filtrate is concentrated, and the residue is purified by column chromatography with hexane-EtOAc giving 221 mg of the product

as a colorless solid.

¹H NMR ([D]₆-DMSO) δ 3.27 (m, 4H), 3.52 (m, 4H), 6.86 (d, 1H), 7.45 (dd, 1H), 7.86 (d, 1H), 8.06 (d, 1H), 8.76 (d, 1H).

5 **7-Chloro-4-[4-(4-fluorobenzylamino(imino)methyl)piperazin-1-yl]quinoline**

7-Chloro-4-(4-cyanopiperazin-1-yl)quinoline (219 mg, 0.8 mmol) and 4-fluorobenzylamine (114 μL, 1.0 mmol) in 1,1,1,3,3,3-hexafluoro-2-propanol (5 mL) are heated at 130 °C in a sealed tube for 44 h. The reaction mixture is concentrated, and 10 the residue is purified by column chromatography with CH₂Cl₂-NH₃-MeOH to give the title product as a yellow foam.

¹H NMR ([D]₆-DMSO) δ 3.17 (m, 4H), 3.49 (m, 4H), 4.17 (s, 2H), 5.80 (br., 2H), 7.01 (d, 1H), 7.09 (t, 2H), 7.35 (dd, 2H), 7.53 (dd, 1H), 7.96 (d, 1H), 8.05 (d, 1H), 8.68 (d, 1H);

15 MS, m/e = 398 (M⁺).

Example 107

Preparation of 7-Chloro-4-[4-(4-fluorophenylamino(imino)methyl)piperazin-1-yl]quinoline

20 7-Chloro-4-(4-cyanopiperazin-1-yl)quinoline (150 mg, 0.55 mmol) and 4-fluoroaniline (66 μL, 0.7 mmol) in 1,1,1,3,3,3-hexafluoro-2-propanol (5 mL) are heated at 150 °C in a sealed tube for 66 h. The reaction mixture is concentrated, and the residue is purified by column chromatography with CH₂Cl₂-NH₃-MeOH to give the title product as 25 a colorless foam.

¹H NMR ([D]₆-DMSO) δ 3.19 (m, 4H), 3.59 (m, 4H), 5.39 (br. s, 2H), 6.70 (dd, 2H), 6.98 (t, 2H), 7.03 (d, 1H), 7.54 (dd, 1H), 7.96 (d, 1H), 8.08 (d, 1H), 8.70 (d, 1H); MS, m/e = 384 (M⁺).

30 **Example 108**

Preparation of 7-Chloro-4-[4-(cycloheptylaminocarbonyl)piperazin-1-yl]quinoline

53001A

As described for example 78, cycloheptylamine (206 mg, 1.82 mmol), 4-nitrophenyl chloroformate (366 mg, 1.82 mmol), diisopropyl(ethyl)amine (554 mg, 4.3 mmol), and 7-chloro-4-(piperazin-1-yl)quinoline (300 mg, 1.21 mmol) are reacted affording the title product after flash chromatography with CH₂Cl₂-MeOH.

5 ¹H NMR (DSMO-d₆/TFA) δ 8.62 (d, 1), 8.21 (d, 1), 7.94 (s, 1), 7.63 (d, 1), 7.16 (d, 1), 6.37 (br, 1), 3.85 (m, 4), 3.61 (m, 1), 3.54 (m, 4), 1.72 (m, 2), 1.24 - 1.61 (m, 10);

Example 109

Preparation of 7-Chloro-4-[4-(cyclooctylaminocarbonyl)piperazin-1-yl]quinoline

10

As described for example 78, cyclooctylamine (234.1 mg, 1.82 mmol), 4-nitrophenyl chloroformate (366 mg, 1.82 mmol), diisopropyl(ethyl)amine (554 mg, 4.3 mmol), and 7-chloro-4-(piperazin-1-yl)quinoline (300 mg, 1.21 mmol) are reacted affording the title product after flash chromatography with CH₂Cl₂-MeOH.

15 ¹H NMR (DSMO-d₆/TFA) δ 8.62 (d, 1), 8.2 (d, 1), 7.97 (s, 1), 7.63 (d, 1), 7.18 (d, 1), 6.32 (br, 1), 3.85 (m, 4), 3.64 (m, 1), 3.57 (m, 4), 1.38 - 1.65 (m, 14);

Example 110

Preparation of 7-Chloro-4-[4-[(4-methylcyclohexyl)aminocarbonyl]piperazin-1-yl]quinoline

20

As described for example 78, 4-methylcyclohexylamine (206.1 mg, 1.82 mmol), 4-nitrophenyl chloroformate (366 mg, 1.82 mmol), diisopropyl(ethyl)amine (554 mg, 4.3 mmol), and 7-chloro-4-(piperazin-1-yl)quinoline (300 mg, 1.21 mmol) are reacted affording the title product after flash chromatography with CH₂Cl₂-MeOH.

25 ¹H NMR (DSMO-d₆/TFA) δ 8.61 (d, 1), 8.2 (d, 1), 7.97 (s, 1), 7.61 (d, 1), 7.19 (d, 1), 3.85 (m, 4), 3.64 (m, 1), 3.37 (m, 4), 1.71 (m, 1), 1.1 -1.67 (m, 9), 0.75 -0.9 (m, 3);

Example 111

Preparation of 7-Chloro-4-[4-[(2-methylcyclohexyl)aminocarbonyl]piperazin-1-yl]quinoline

As described for example 78, 2-methylcyclohexylamine (170 mg, 1.5 mmol), 4-nitrophenyl chloroformate (302 mg, 1.5 mmol), diisopropyl(ethyl)amine (469 mg, 3.7 mmol), and 7-chloro-4-(piperazin-1-yl)quinoline (248 mg, 1.0 mmol) are reacted affording the title product after flash chromatography with CH₂Cl₂-MeOH.

5 ¹H NMR (DSMO-d₆/TFA) δ 8.61 (d, 1), 8.2 (d, 1), 8.0 (s, 1), 7.62 (d, 1), 7.17 (d, 1), 3.91 (m, 4), 3.75 (m, 1), 3.58 (m, 4), 0.9 -1.8 (m, 9), 0.86 (m, 3);

Example 112

Preparation of 2-(2-Aminoethylamino)-7-chloro-4-[4-(4-

10 **fluorophenylaminocarbonyl)piperazin-1-yl]quinoline**

To 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-[2-(trifluoroacetamino)ethylamino]quinoline (13 mg, 0.024 mmol) was added THF (0.5 mL) and 0.5 N KOH (2.0 eq, 0.14 mL), the mixture was stirred at rt. for 2h, additional 15 0.5 N KOH (0.5 mL, 10eq.) was added and the reaction mixture was heated at 50°C for 30 min, MeOH (0.2 ml) was added and heated at the same temperature for 30 min,. Evaporation and dilution with AcOEt, washing with brine, usual workup gave the residue, which was purified by preparative TLC with 20% MeOH in DCM with 0.5% hydroxylamine by twice developing the plate to afford the title compound.

20 ¹H NMR (CD₃OD) δ 3.10 (m, 4H), 3.25 (m, 2H), 3.50 (m, 2H), 3.75 (m, 4H), 6.5 (s., 1H), 6.95 (m, 2H), 7.10 (m, 1H), 7.30 (m, 2H), 7.50 (s, 1H), 7.80 (m, 1H).

Example 113

Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-[2-

25 **(trifluoroacetylamino)ethylamino]quinoline**

2-[[(Trifluoroacetylamino)ethyl]amino]-7-chloro-4-[4-(t-butoxycarbonyl)piperazin-1-yl]quinoline (31 mg, 0.062 mmol) was treated with TFA:DCM (1:1, 0.2 mL) at r. t. for 40 min. Evaporation of the solvents, dilution with DCM and evaporation afford the 30 corresponding unsubstituted piperazine as the TFA salt which was dissolved in DCM (15 mL) cooled in an ice bath. NEt₃ (79 µl, 9 eq.) and 4-fluorophenyl isocyanate (8 µl, 1.5 eq.) was added. After stirring at rt. for 1h, the mixture was diluted with DCM and

washed with water followed by the usual work up. Flash chromatography on silica gel with MeOH in DCM afforded the title compound.

¹H NMR (CD₃OD) δ 3.0 (m, 4H), 3.4 (m, 2H), 3.55 (m, 2H), 3.70 (m, 4H), 6.2 (s., 1H), 6.95 (m, 2H), 7.05 (d, 1H), 7.30 (m, 2H), 7.50 (s, 1H), 7.75 (d, 1H).

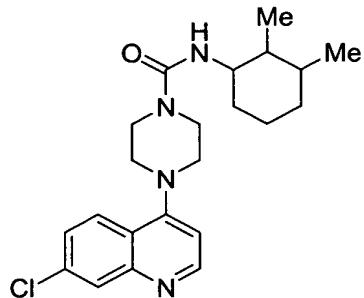
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Example 114

Preparation of 7-Chloro-4-[4-(4-fluorophenylaminocarbonyl)piperazin-1-yl]-2-methoxyquinoline

To a solution of 2,4,7-trichloroquinoline (150 mg, 0.685 mmol) in toluene (3 mL) was added a suspension of NaOMe (136 mg, 3.4 eq.) in toluene (3 mL). The mixture was refluxed at 130°C for 24 h. After cooling to r. t. the solid was filtered and the filtrate was evaporated. The residue was dissolved in MeOH and evaporated to afford 2-methoxy-4,7-dichloroquinoline (139 mg) as an off-white solid. To it was added 1-N-Boc piperazine (455 mg, 4.0 eq.). The mixture was heated in a sealed tube at 150°C for 3h, additional piperazine (6 eq.) was added and treated with n-BuOH (2 mL). The reaction mixture was heated at the same temperature overnight. Evaporation of solvent under vacuum and flash chromatography on silica gel with AcOEt in hexane affords the corresponding t-butoxy-carbonylpiperazinylquinoline (20 mg), which was treated with TFA:DCM(1:1, 0.4 mL) at rt. for 1h . Evaporation of the solvents and dilution with DCM and evaporation affords the corresponding unsubstituted piperazine as the TFA salt, which was dissolved in DCM (15 mL), in an ice bath. NEt₃ (7 eq.) and 4-fluorophenylisocyanate (9μl, 1.5 eq.) was added. After stirring at rt. for 3h, the mixture was diluted with DCM and washed with water, followed by the usual work up. Flash chromatography on silica gel with MeOH in DCM afforded the title compound.

¹H NMR (CDCl₃) δ 3.20 (m, 4H), 3.75 (m, 4H), 4.05 (s., 3H), 6.35 (s, 1H), 6.45 (s, 1H), 7.00 (t, 2H), 7.30- (m, 3H), 7.80 (s, 1H), 7.85 (d, 1H).

Example 115 :**Preparation of 4-(7-Chloro-4-quinolinyl)-N-(2,3-dimethylcyclohexyl)-1-piperazinecarboxamide**

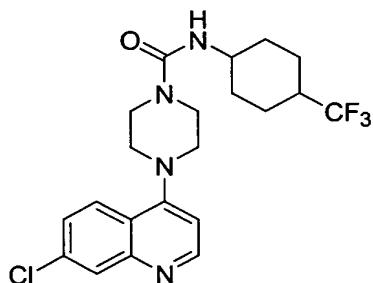
5

As described for example 78, 2,3-dimethyl-cyclohexanamine (154 mg, 1.21 mmol), 4-nitrophenyl chloroformate (244 mg, 1.21 mmol), diisopropyl(ethyl)amine (209 mg, 1.62 mmol), and 7-chloro-4-(piperazin-1-yl)quinoline (200 mg, 0.81 mmol) are reacted affording the title product after flash chromatography with CH₂Cl₂-MeOH.

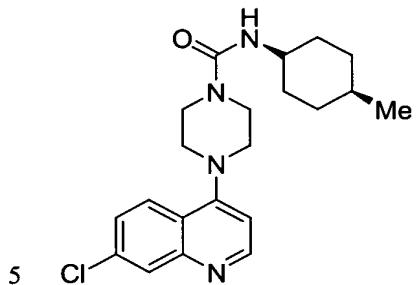
¹⁰ ^{1H} NMR (DSMO-d₆) δ 0.7-0.9 (m, 7H), 1.13-1.92 (m, 8H), 3.2 (m, 1H), 3.6 (br. s, 4H), 3.9 (br. s, 4H), 7.2 (m, 1H), 7.7 (m, 1H), 8.0 (s, 1H), 8.2 (m, 1H), 8.6 (m, 1H).

Example 116:**Preparation of 4-(7-Chloro-4-quinolinyl)-N-[4-(trifluoromethyl)cyclohexyl]-1-piperazinecarboxamide**

¹⁵



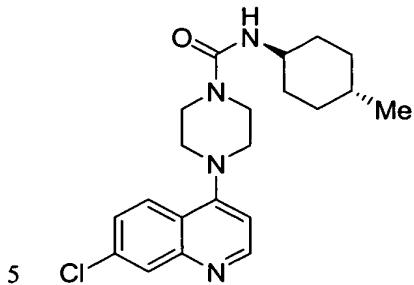
As described for example 78, 4-trifluoromethyl cyclohexyl amine, 4-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(piperazin-1-yl)quinoline are reacted to afford the product after flash chromatography with CH₂Cl₂-MeOH. LC-MS: 441 (M⁺+1). ^{1H} NMR (CDCl₃) δ 8.78 (d, 1H), 8.05 (s, 1H), 7.95 (d, 1H), 7.46 (d, 1H), 6.85 (d, 1H), 4.60 (d, 1H), 4.03 (m, 1H), 3.65 (m, 4H), 3.20 (m, 4H), 2.10 (m, 1H), 1.85 (m, 4H), 1.50-1.70 (m, 4H).

Example 117:**Preparation of 4-(7-Chloro-4-quinolinyl)-N-(*cis*-4-methylcyclohexyl)-1-piperazinecarboxamide**

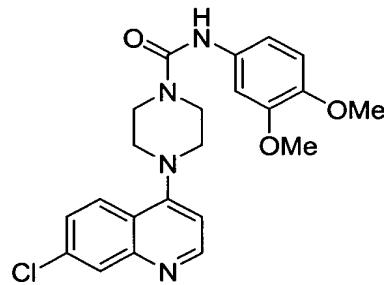
Preparation of cis-4-methyl-1-cyclohexanamine: To a solution of trans-4-methyl-1-cyclohexanol (5 g, 44 mmol) and Et₃N (8.9 g, 88 mmol) in CH₂Cl₂ at 0 °C was added MsCl (6.03 g, 53 mmol). After 2 h at 0 °C, the reaction was quenched with NaHCO₃ (sat.), extracted with EtOAc, and dried over Na₂SO₄. Concentration in vacuo afforded 8.9 g of trans-4-methyl-1-methanesulfonatecyclohexanol. The mixture of trans-4-methyl-1-methanesulfonatecyclohexanol (8.9 g) with NaN₃ (28.5 g, 0.44 mol) in DMSO (300 mL) was kept at 95 °C for 6 h, then cooled to rt, and poured into ice-water (300 mL). The reaction mixture was extracted with EtOAc (3x200 mL), washed with brine, and dried over Na₂SO₄. Concentration in vacuo afforded crude *cis*-1-(4-methylcyclohexyl)azide. To a suspension of LiAlH₄ (2.5 g, 66 mmol) in THF (100 mL) was added *cis*-1-(4-methylcyclohexyl)azide in THF (15 mL) at 0 °C. After 12 h at rt, the reaction was quenched with 6 N NaOH (20 mL), and the solid was filtered off. The resulting filtrate was concentrated in vacuo to give 7.2 g of *cis*-4-methyl-1-cyclohexanamine.

20

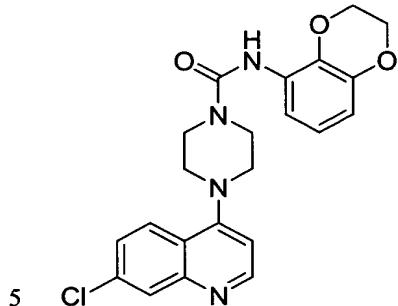
*Preparation of 4-(7-chloro-4-quinolinyl)-N-(*cis*-4-methylcyclohexyl)-1-piperazinecarboxamide:* As described for example 78, *cis*-4-methyl-1-cyclohexanamine (170 mg, 1.5 mmol), 4-nitrophenyl chloroformate (302 mg, 1.5 mmol), diisopropyl(ethyl)amine (258 mg, 2 mmol), and 7-chloro-4-(piperazin-1-yl)quinoline (248 mg, 1 mmol) are reacted affording the title product after flash chromatography with CH₂Cl₂-MeOH. ¹H NMR (DSMO-d₆) δ 0.92 (d, 3H), 1.2 (m, 2H), 1.55-1.7 (m, 8H), 3.22 (m, 4H), 3.65 (m, 4H), 4.02 (br. s, 1H), 4.6 (m, 1H), 6.82 (d, 1H), 7.44 (d, 1H), 7.98 (d, 1H), 8.06 (s, 1), 8.78 (d, 1H).

Example 118 :**Preparation of 4-(7-Chloro-4-quinolinyl)-N-(*trans*-4-methylcyclohexyl)-1-piperazin carboxamid**

As described for example 78, *trans*-4-methyl-1-cyclohexanamine (170 mg, 1.5 mmol) prepared in a similar manner as for *cis*-4-methyl-1-cyclohexanamine, 4-nitrophenyl chloroformate (302 mg, 1.5 mmol), diisopropyl(ethyl)amine (258 mg, 2 mmol), and 7-chloro-4-(piperazin-1-yl)quinoline (248 mg, 1 mmol) are reacted affording the title product after flash chromatography with CH₂Cl₂-MeOH. ¹H NMR (DSMO-d₆) δ 0.9 (d, 3H), 1.0-1.2 (m, 3H), 1.23-1.4 (m, 2H), 1.7 (m, 2H), 2.02 (m, 2H), 3.2 (m, 4H), 3.61 (m, 4H), 4.38 (m, 1H), 6.83 (d, 1H), 7.43 (d, 1H), 7.94 (d, 1H), 8.05 (s, 1H), 8.77 (d, 1H).

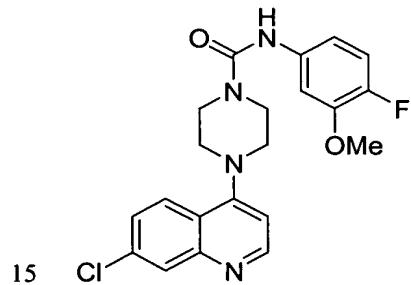
Example 119:**15 Preparation of 4-(7-chloro-4-quinolinyl)-N-(3,4-dimethoxyphenyl)- 1-piperazinecarboxamide**

As described for example 78, 3,4-dimethoxyaniline, *p*-nitrophenyl chloroformate, diisopropylethyl amine, and 7-chloro-4-piperazinylquinoline are reacted to afford the product. LC-MS: 427 (M⁺+1). ¹H NMR (CDCl₃) δ 8.80 (d, 1H), 8.05 (s, 1H), 7.95 (d, 1H), 7.50 (d, 1H), 7.20 (s, 1H), 6.85 (d, 1H), 6.80 (d, 1H), 6.75 (m, 1H), 6.40 (s, 1H), 3.90 (s, 3H), 3.85(s, 3H), 380 (m, 4H), 3.25 (m, 4H).

Example 120:**Preparation of 4-(7-Chloro-4-quinolinyl)-N-(2,3-dihydro-1,4-benzodioxin-6-yl)-1-piperazinecarboxamide**

As described for example 78, 5-amino-1,4-benzodioxane, *p*-nitrophenyl chloroformate, diisopropylethyl amine, and 7-chloro-4-piperazinylquinoline are reacted to afford the product. LC-MS: 425 (M^++1). 1H NMR (DMSO- d_6) δ 8.70 (d, 1H), 8.50 (s, 1H), 8.05 (d, 1H), 7.95 (s, 1H), 7.55 (d, 1H), 7.00 (m, 2H), 6.85 (d, 1H), 6.70 (d, 1H), 4.15 (m, 4H), 3.65 (m, 4H), 3.20 (m, 4H).

10

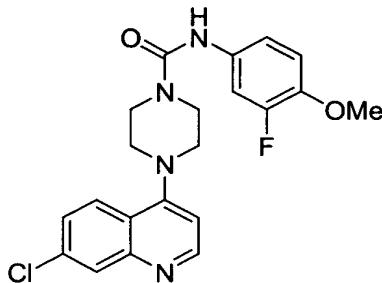
Example 121 :**Preparation of 4-(7-Chloro-4-quinolinyl)-N-(4-fluoro-3-methoxyphenyl)-1-piperazinecarboxamide**

As described for example 78, 3-methoxy-4-fluoro-aniline, *p*-nitrophenyl chloroformate, diisopropylethyl amine, and 7-chloro-4-piperazinylquinoline are reacted to afford the product. LC-MS: 415 (M^++1). 1H NMR (CDCl₃) δ 8.78 (d, 1H), 8.05 (s, 1H), 7.95 (d, 1H), 7.50 (d, 1H), 7.35 (d, 1H), 7.00 (m, 1H), 6.85 (d, 1H), 6.70 (m, 1H), 6.65 (s, 1H), 3.85 (s, 3H), 3.75 (m, 4H), 3.25 (m, 4H).

20

Example 122 :

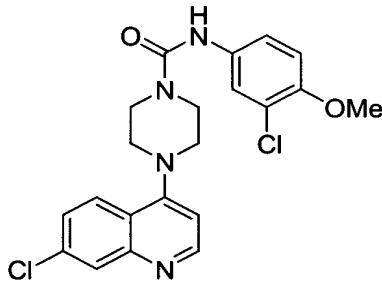
Preparation of 4-(7-Chloro-4-quinolinyl)-*N*-(3-fluoro-4-methoxyphenyl)-1-piperazinecarboxamide



As described for example 78, 4-methoxy-3-fluoro-aniline, *p*-nitrophenyl chloroformate, diisopropylethyl amine, and 7-chloro-4-piperazinylquinoline are reacted to afford the product. LC-MS: 415 (M^++1). ^1H NMR (CDCl_3) δ 8.78 (d, 1H), 8.05 (s, 1H), 7.95 (d, 1H), 7.50 (d, 1H), 7.35 (d, 1H), 7.00 (m, 1H), 6.85 (d, 1H), 6.70 (m, 1H), 6.60 (s, 1H), 3.85 (s, 3H), 3.78 (m, 4H), 3.25 (m, 4H).

10 **Example 123 :**

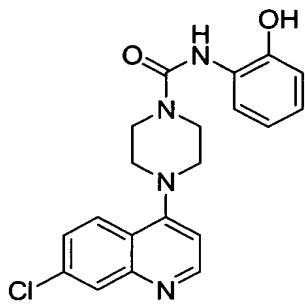
Preparation of *N*-(3-Chloro-4-methoxyphenyl)-4-(7-chloro-4-quinolinyl)-1-piperazinecarboxamide



As described for example 78, 4-methoxy-3-chloro-aniline, *p*-nitrophenyl chloroformate, diisopropylethyl amine, and 7-chloro-4-piperazinylquinoline are reacted to afford the product. LC-MS: 431 (M^++1). ^1H NMR (CDCl_3) δ 8.78 (d, 1H), 8.05 (s, 1H), 7.95 (d, 1H), 7.50 (d, 1H), 7.45 (s, 1H), 7.22 (d, 1H), 6.90 (m, 2H), 6.38 (s, 1H), 3.85 (s, 3H), 3.80 (m, 4H), 3.25 (m, 4H).

20 **Example 124 :**

Preparation of 4-(7-Chloro-4-quinolinyl)-*N*-(2-hydroxyphenyl)-1-piperazinecarboxamide



Preparation of 4-(7-chloro-4-quinolinyl)-N-(2-methoxyphenyl)-1-

piperazinecarboxamide: As described for example 78, 2-methoxyaniline, 4-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(piperazin-1-yl)quinoline are

5 reacted to afford the product after flash chromatography with CH₂Cl₂-MeOH.

Preparation of 4-(7-chloro-4-quinolinyl)-N-(2-hydroxyphenyl)-1-piperazinecarboxamide:

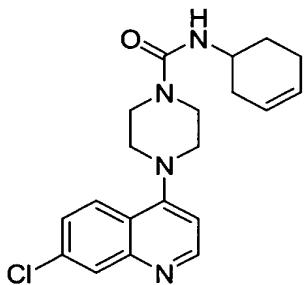
To a stirred solution of 4-(7-chloro-4-quinolinyl)-N-(2-methoxyphenyl)-1-piperazine carboxamide (40 mg, 0.1 mmol) in CH₂Cl₂ (6 mL) was added a solution of BBr₃ (0.2

10 mL, 1.0 M) at rt. The reaction mixture was stirred at rt overnight, then quenched with 1N NaOH solution. The reaction mixture was extracted with CH₂Cl₂, washed with brine, and dried over Na₂SO₄. Concentration in vacuo followed by flash chromatography afforded the title product. LC-MS 383 (M⁺+1). ¹H NMR (CDCl₃) δ 8.75 (d, 1H), 8.05 (s, 1H), 7.95 (d, 1H), 7.45 (d, 1H), 7.05 (m, 3H), 6.85 (m, 2H), 6.70 (s, 1H), 3.85 (m, 4H), 3.30 (m, 4H).

Example 125 :

Preparation of 4-(7-Chloro-4-quinolinyl)-N-(3-cyclohexen-1-yl)-1-

piperazinecarboxamide

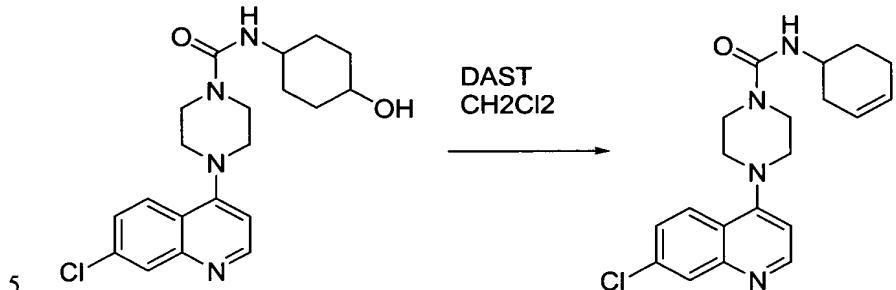


20

Preparation of 4-(7-chloro-4-quinolinyl)-N-(4-hydroxycyclohexyl)-1-

piperazinecarboxamide: As described for example 78, 4-aminocyclohexanol (84 mg, 0.73 mmol), 4-nitrophenyl chloroformate (146 mg, 0.73 mmol), diisopropyl(ethyl)amine

(320 mg, 2.4 mmol), and 7-chloro-4-(piperazin-1-yl)quinoline (150 mg, 0.81 mmol) are reacted affording 80 mg of the product after flash chromatography with CH₂Cl₂-MeOH. ¹H NMR (DSMO-d₆) δ 1.17 (m, 4H), 1.74 (m, 4H), 3.38 (m, 2H), 3.56 (br. s, 4H), 3.82 (br., s, 4H), 7.16 (d, 1H), 7.64 (dd, 1H), 1.97 (d, 1H), 8.2 (d, 1H), 8.62 (d, 1H).



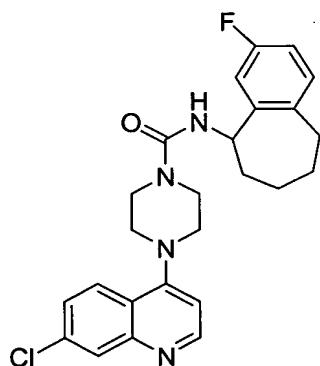
Preparation of 4-(7-chloro-4-quinolinyl)-N-(3-cyclohexen-1-yl)-1-piperazinecarboxamide: To a stirred solution of 4-(7-chloro-4-quinolinyl)-N-(4-hydroxycyclohexyl)-1-piperazinecarboxamide (100 mg) in CH₂Cl₂ at -78 °C was added DAST (100 mg). The reaction was warmed to rt overnight, and quenched with NaOH (sat.). The reaction mixture was extracted with EtOAc, and dried. Concentration in vacuo followed by flash chromatography with CH₂Cl₂-MeOH afforded the title product.

10 ¹H NMR (DSMO-d₆) δ 1.03 (m, 1H), 1.4 (m, 1H), 1.7 (m, 1H), 1.9 (m, 1H), 2.05 (m, 1H), 2.1 (m, 1H), 3.1 (br. s, 4H), 3.6 (br. s, 4H), 3.65 (m, 1H), 5.6 (m, 2H), 6.4 (d, 1H), 7.0 (d, 1H), 7.6 (m, 1H), 8.0 (s, 1H), 8.1 (d, 1H), 8.7 (d, 1H).

15

Example 126 :

Preparation of 4-(7-Chloro-4-quinolinyl)-N-(3-fluoro-6,7,8,9-tetrahydro-5H-benzo[α]cyclohepten-5-yl)-1-piperazinecarboxamide

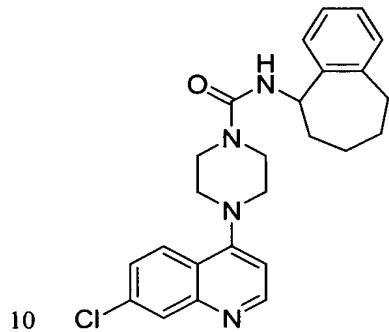


20 As described for example 78, 8-fluoro-benzocycloheptyl-1-amine hydrochloride, *p*-nitrophenyl chloroformate, diisopropylethyl amine, and 7-chloro-4-piperazinylquinoline

are reacted to afford the product after flash chromatography with CH₂Cl₂-MeOH. LC-MS: 453 (M⁺+1). ¹H NMR (CDCl₃) δ 8.76 (d, 1H), 8.05 (s, 1H), 7.96 (d, 1H), 7.45 (d, 1H), 7.05 (m, 1H), 6.95 (m, 1H), 6.85 (d, 1H), 6.80 (m, 1H), 5.12 (m, 1H), 4.85 (m, 1H), 3.70 (m, 4H), 3.25 (m, 4H), 2.85 (m, 2H), 1.95 (m, 2H), 1.80 (m, 2H), 1.67 (m, 1H), 5 1.50 (m, 1H).

Example 127 :

Preparation of 4-(7-Chloro-4-quinolinyl)-N-(6,7,8,9-tetrahydro-5H-benzo[*a*]cyclohepten-5-yl)-1-piperazinecarboxamide

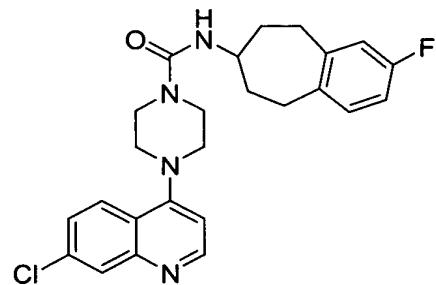


10

As described for example 78, benzocycloheptyl-1-amine hydrochloride, *p*-nitrophenyl chloroformate, diisopropylethyl amine, and 7-chloro-4-piperazinylquinoline are reacted to afford the product. LC-MS: 435 (M⁺+1). ¹H NMR (DMSO-d₆) δ 8.60 (s, 1H), 8.05 (d, 1H), 7.90 (s, 1H), 7.50 (d, 1H), 7.15 (m, 1H), 7.00 (m, 4H), 4.95 (m, 1H), 3.65 (m, 4H), 15 3.18 (m, 4H), 2.85 (m, 2H), 1.85 (m, 3H), 1.70 (m, 1H), 1.58 (m, 1H), 1.20 (m, 1H).

Example 128 :

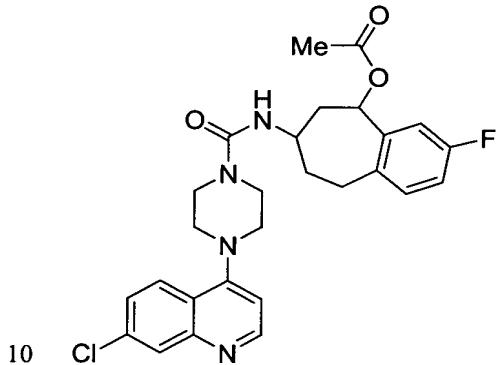
Preparation of 4-(7-Chloro-4-quinolinyl)-N-(2-fluoro-6,7,8,9-tetrahydro-5H-benzo[*a*]cyclohepten-7-yl)-1-piperazinecarboxamide



As described for example 78, 8-fluoro-benzocyloheptyl-3-amine, *p*-nitrophenyl chloroformate, diisopropylethyl amine, and 7-chloro-4-piperazinylquinoline are reacted to afford the product. LC-MS: 453 (M^++1). ^1H NMR (CD_3OD) δ 8.50 (d, 1H), 8.25 (d, 1H), 7.90 (s, 1H), 7.65 (d, 1H), 7.15 (d, 1H), 7.05 (m, 1H), 6.85 (m, 1H), 6.78 (m, 1H),
5 3.95 (m, 4H), 3.90 (m, 1H), 3.70 (m, 4H), 2.78 (m, 4H), 2.10 (m, 2H), 1.35 (m, 2H).

Example 129 :

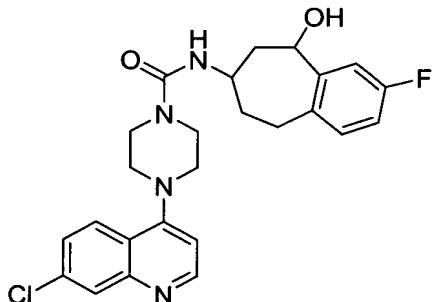
Preparation of 4-(7-Chloro-4-quinolinyl)- *N*-[9-(acetyloxy)-2-fluoro-6,7,8,9-tetrahydro-5*H*-benzo[α]cyclohepten-7-yl]-1-piperazinecarboxamide



As described for example 78, 8-fluoro-benzocyloheptyl-1-acetyloxy-3-amine, *p*-nitrophenyl chloroformate, diisopropylethyl amine and 7-chloro-4-piperazinylquinoline are reacted to afford the product. LC-MS: 512 (M^++1). ^1H NMR (CD_3OD) δ 8.50 (d, 1H), 8.25 (d, 1H), 7.90 (s, 1H), 7.65 (d, 1H), 7.15 (d, 2H), 7.05 (m, 1H), 6.85 (m, 1H),
10 5.90 (m, 1H), 4.10 (m, 1H), 3.95 (m, 4H), 3.75 (m, 4H), 2.80 (m, 2H), 2.26 (m, 1H),
15 2.20 (s, 3H), 2.15 (m, 1H), 1.65 (m, 1H), 1.30 (m, 1H).

Example 130 :

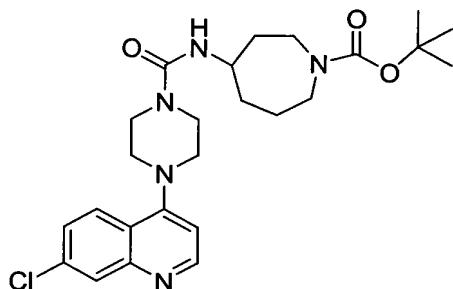
Preparation of 4-(7-Chloro-4-quinolinyl)-*N*-(2-fluoro-6,7,8,9-tetrahydro-9-hydroxy-5*H*-benzo[α]cyclohepten-7-yl)-1-piperazinecarboxamide



To a stirred solution of N-[9-(acetyloxy)-2-fluoro-6,7,8,9-tetrahydro-5H-benzo[a]cyclohepten-7-yl]-4-(7-chloro-4-quinolinyl)-1-piperazinecarboxamide (30.0 mg, 0.0588 mmol) in THF:H₂O (3:1, 2 mL) was added LiOH (8 mg). The reaction mixture
5 was stirred at rt for 3h, then THF was evaporated in vacuo. The reaction mixture was diluted with AcOEt (100 mL), washed with brine, and dried. Concentration followed by flash chromatography afforded the title product. LC- MS: 469 (M⁺+1). δ ¹H NMR (CDCl₃) δ 8.70 (d, 1H), 8.05 (s, 1H), 7.95 (d, 1H), 7.45 (d, 1H), 7.05 (m, 2H), 6.85 (m, 2H), 4.95 (m, 1H), 4.35 (m, 1H), 3.65 (m, 4H), 3.20 (m, 4H), 3.15 (m, 1H), 2.60 (m, 10 1H), 2.20 (m, 2H), 1.8 (m, 3H).

Example 131 :

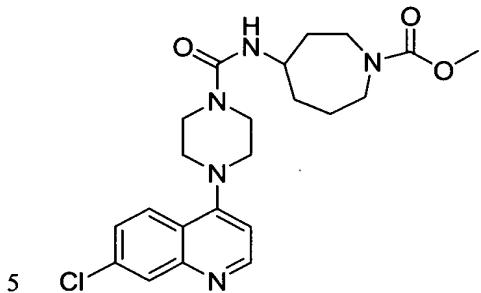
Preparation of 4-[[[4-(7-Chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-1*H*-azepine-1-carboxylic acid, 1,1-dimethylethyl ester



As described for example 78, 4-amino-N-1-Boc-azepine, *p*-nitrophenyl chloroformate, diisopropylethyl amine and 7-chloro-4-piperazinylquinoline are reacted to afford the title product. LC-MS: 488 (M⁺+1). δ ¹H NMR (CDCl₃) δ 8.75 (d, 1H), 8.05 (s, 1H), 7.96 (d, 1H), 7.45 (d, 1H), 6.85 (d, 1H), 4.55 (m, 1H), 4.05 (m, 1H), 3.90 (m, 1H), 3.60 (m, 4H), 3.30 (m, 1H), 3.20 (m, 4H), 3.10 (m, 2H), 2.10 (m, 1H), 1.60-1.90 (m, 3H), 1.50 (s, 9H).

Example 132 :

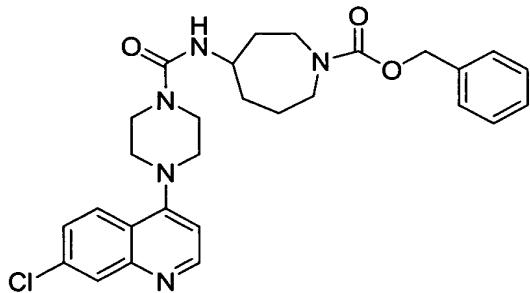
Preparation of 4-[[[4-(7-Chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-1*H*-azepine-1-carboxylic acid, methyl ester



To a mixture of 4-(7-chloro-4-quinolinyl)-*N*-(hexahydro-1*H*-azepin-4-yl)-1-piperazine carboxamide (40 mg, 0.082 mmol) obtained by the de-protection of 4-[[[4-(7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-1*H*-azepine-1-carboxylic acid 1,1-dimethylethyl ester with TFA and Cs₂CO₃ (40 mg) in DMF (3 mL), MeI (11.6 mg) was introduced. The reaction mixture was stirred at 50 °C for 30 min, then MeI (30 mg.) was added. After 1h at rt, the reaction mixture was diluted with water and extracted with AcOEt, washed with water and brine, and dried. Concentration *in vacuo* followed by flash chromatography afforded the title product. LC-MS: 446 (M⁺+1). ¹H NMR (CDCl₃) δ 8.78 (d, 1H), 8.05 (s, 1H), 7.95 (d, 1H), 7.45 (d, 1H), 6.85 (d, 1H), 4.45 (m, 1H), 3.90 (m, 1H), 3.70 (s, 3H), 3.65 (m, 4H), 3.20 (m, 5H), 2.10 (m, 1H), 1.95 (m, 2H), 1.60-1.70 (m, 3H), 1.27 (m, 2H).

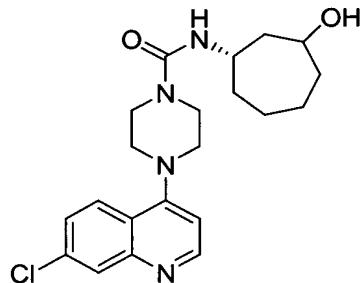
Example 133 :

20 **Preparation of 4-[[[4-(7-Chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-1*H*-azepine-1-carboxylic acid, phenylmethyl ester**



As described for example 132, 4-(7-chloro-4-quinolinyl)-N-(hexahydro-1H-azepin-4-yl)-1-piperazinecarboxamide obtained from the de-protection of 4-[[[4-(7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-1H-azepine-1-carboxylic acid 1,1-dimethylethyl ester with TFA, Cs₂CO₃, and CbzCl are reacted to afford the title product. LC-MS: 522 (M⁺+1), ¹H NMR (CDCl₃) δ 8.75 (d, 1H), 8.05 (s, 1H), 7.95 (d, 1H), 7.45 (d, 1H), 7.30 (m, 5H), 6.80 (d, 1H), 5.15 (m, 2H), 4.50 (m, 1H), 3.65-3.95 (m, 3H), 3.58 (m, 4H), 3.30 (m, 1H), 3.18 (m, 4H), 2.10 (m, 1H), 1.65-1.95 (m, 3H), 1.60 (m, 2H).

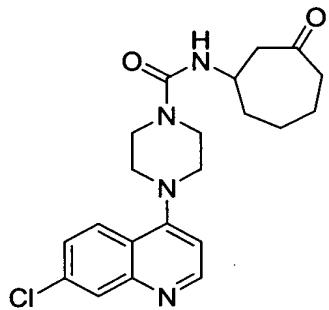
10

Example 134 :**Preparation of 4-(7-Chloro-4-quinolinyl)-N-(3-hydroxycycloheptyl)-1-piperazinecarboxamide**

15 As described for example 78, 3-hydroxycycloheptyl-amine, p-nitrophenyl chloroformate, diisopropylethyl amine and 7-chloro-4-piperazinylquinoline are reacted to afford the product. LC-MS: 403 (M⁺+1). ¹H NMR (CDCl₃) δ 8.75 (d, 1H), 8.05 (s, 1H), 7.96 (d, 1H), 7.45 (d, 1H), 6.84 (d, 1H), 4.50 (d, 1H), 4.10 (m, 2H), 3.65 (m, 4H), 3.20 (m, 4H), 2.38 (m, 1H), 2.0 (m, 3H), 1.80 (m, 4H), 1.40 (m, 2H).

20

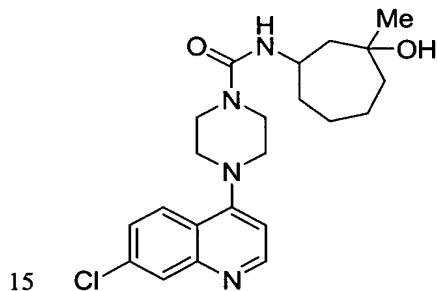
Example 135 :**Preparation of 4-(7-Chloro-4-quinolinyl)-N-(3-oxocycloheptyl)-1-piperazinecarboxamide**



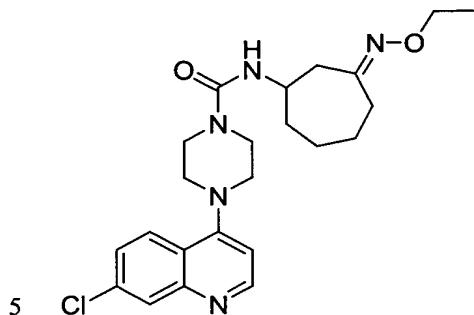
To a stirred solution of 4-(7-chloro-4-quinolinyl)-N-(3-hydroxycycloheptyl)-1-piperazinecarboxamide (300 mg, 0.744 mmol) in CH_2Cl_2 (20 mL), the Dess-Martin reagent (348 mg, 0.82 mmol) was added at rt. After 3 h, the reaction mixture was quenched with 1g of $\text{Na}_2\text{S}_2\text{O}_3$ and 5 mL of NaHCO_3 (sat.). The reaction mixture was extracted with CH_2Cl_2 , washed with brine, and dried over Na_2SO_4 . Concentration *in vacuo* followed by flash chromatography afforded the title product. LC-MS: 401 (M^++1).
 5 ^1H NMR (CDCl_3) δ 8.78 (d, 1H), 8.05 (s, 1H), 7.95 (d, 1H), 7.45 (d, 1H), 6.85 (d, 1H), 4.70 (m, 1H), 4.25 (m, 1H), 3.60 (m, 4H), 3.25 (m, 4H), 3.00 (m, 1H), 2.45-2.65 (m,
 10 3H), 1.70-1.90 (m, 6H).

Example 136 :

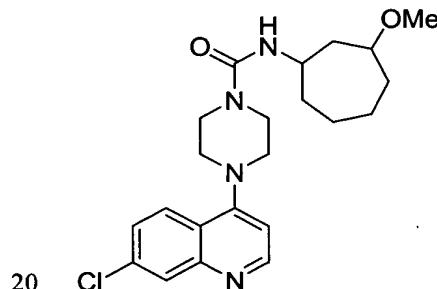
Preparation of 4-(7-Chloro-4-quinolinyl)-N-(3-hydroxy-3-methylcycloheptyl)-1-piperazinecarboxamide



To a stirred solution of 4-(7-chloro-4-quinolinyl)-N-(3-oxocycloheptyl)-1-piperazinecarboxamide (8 mg, 0.02 mmol) in THF (3 mL) was added a solution of MeMgBr (3M, 0.1 mL) at 0 °C. After 5 h at 0 °C, the reaction mixture was quenched with MeOH and Na_2CO_3 (sat. 50 mL). The reaction mixture was extracted with CH_2Cl_2 , washed with brine, and dried over Na_2SO_4 . Concentration *in vacuo* followed by preparative TLC gave the title product. LC-MS: 401 (M^++1). ^1H NMR (CDCl_3) δ 8.78 (d, 1H), 8.05 (s, 1H), 7.95 (d, 1H), 7.45 (m, 1H), 6.85 (d, 1H), 5.75 (d, 1H), 4.45 (d, 1H), 4.05 (m, 1H), 3.65 (m, 4H), 3.20 (m, 4H), 1.55-1.95 (m, 10H), 1.35 (s, 3H).
 15
 20

Example 137 :**Preparation f 4-(7-Chloro-4-quinolinyl)-N-[(3*E*)-3-(thoxyimino)cycloheptyl]-1-piperazinecarboxamid**

To a stirred solution of 4-(7-chloro-4-quinolinyl)-N-(3-oxocycloheptyl)-1-piperazinecarboxamide (30 mg, 0.075 mmol) in EtOH (3 mL) was added ethoxyamine hydrochloride (36 mg, 0.225 mmol), followed by NaOAc (31 mg, 0.225 mmol). The mixture was heated at reflux for 2h. After cooling to rt , the reaction mixture was poured into water (100 mL), extracted with AcOEt, washed with brine, and dried over Na₂SO₄. Concentration in vacuo followed by flash chromatography afforded the title product. LC-MS: 444 (M⁺+1). ¹H NMR (CDCl₃) δ 8.78 (d, 1H), 8.05 (s, 1H), 7.95 (d, 1H), 7.45 (m, 1H), 6.85 (d, 1H), 4.80 (d, 1H), 4.10 (m, 3H), 3.6 (m, 4H), 3.20 (m, 4H), 2.30-2.90 (m, 4H), 1.50-1.95 (m, 6H), 1.25 (t, 3H).

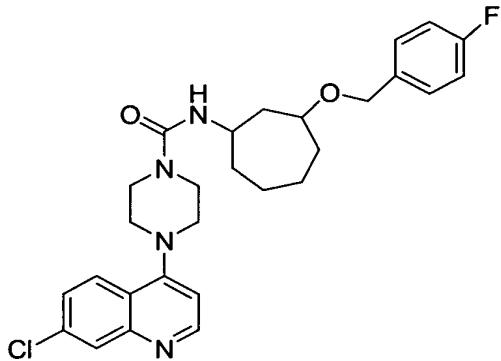
Example 138 :**Preparation of 4-(7-Chloro-4-quinolinyl)-N-(3-methoxycycloheptyl)-1-piperazinecarboxamide**

To a stirred solution of 4-(7-chloro-4-quinolinyl)-N-(3-hydroxycycloheptyl)-1-piperazinecarboxamide (30 mg, 0.074 mmol) in DMF (3 mL) was added NaH (8.9 gm,

0.37 mmol) at rt. After 30 min, MeI (11.5 mg, 0.09 mmol) was added, and the reaction mixture was heated at rt for 2h. After cooling to rt, the reaction mixture was quenched with MeOH, then poured into water (100 mL). The reaction mixture was extracted with EtOAc, washed with brine, and dried over Na₂SO₄. Concentration *in vacuo* followed by 5 flash chromatography afforded the title product. LC-MS: 416 (M⁺+1). ¹H NMR (CDCl₃) δ 8.75 (d, 1H), 8.05 (s, 1H), 7.95 (d, 1H), 7.45 (d, 1H), 6.85 (d, 1H), 4.40 (d, 1H), 4.10 (m, 1H), 3.65 (m, 4H), 3.45 (m, 1H), 3.36 (s, 3H), 3.20 (m, 4H), 2.15 (m, 1H), 2.05 (m, 1H), 1.50-1.90 (m, 6H), 1.40 (m, 2H).

10 **Example 139 :**

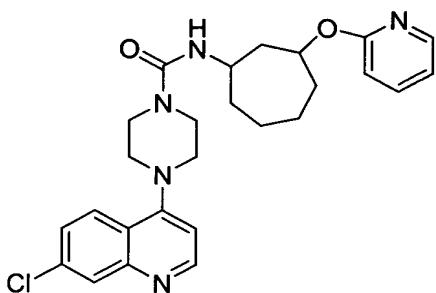
Preparation of 4-(7-Chloro-4-quinolinyl)-N-[3-[(4-fluorophenyl)methoxy]cycloheptyl]-1-piperazinecarboxamide



15 As described for example 138, 4-(7-chloro-4-quinolinyl)-N-(3-hydroxycycloheptyl)-1-piperazinecarboxamide (50 mg, 0.124 mmol), NaH (8.9 mg, 0.37 mmol), and 4-fluorobenzyl bromide (28.2 mg, 0.15 mmol) are reacted to give the title product after 20 flash chromatography with CH₂Cl₂-MeOH. LC-MS: 511 (M⁺+1). ¹H NMR (CDCl₃) δ 8.75 (d, 1H), 8.05 (s, 1H), 7.95 (d, 1H), 7.45 (d, 1H), 7.35 (m, 2H), 7.00 (m, 2H), 6.85 (d, 1H), 4.50 (m, 2H), 4.40 (m, 1H), 4.10 (m, 1H), 3.65 (m, 4H), 3.20 (m, 4H), 2.20 (m, 1H), 2.10 (m, 1H), 1.50-1.90 (m, 6H), 1.40 (m, 2H).

Example 140 :

Preparation of 4-(7-Chloro-4-quinolinyl)-N-[3-(2-pyridinyloxy)cycloheptyl]-1-piperazinecarboxamide

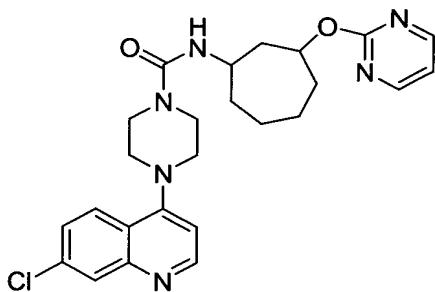


As described for example 138, 4-(7-chloro-4-quinolinyl)-N-(3-hydroxycycloheptyl)-1-piperazinecarboxamide, NaH, and 2-fluoropyridine are reacted to give the product

5 after flash chromatography with CH₂Cl₂-MeOH. LC-MS: 480 (M⁺+1). ¹H NMR (CDCl₃) δ 8.75 (d, 1H), 8.05 (s, 1H), 8.0 (m, 2H), 7.60 (m, 1H), 7.45 (d, 1H), 6.85 (m, 2H), 6.75 (d, 1H), 6.05 (d, 1H), 5.20 (m, 1H), 4.10 (m, 1H), 3.75 (m, 4H), 3.25 (m, 4H), 2.28 (m, 1H), 2.10 (m, 3H), 1.85 (m, 2H), 1.70 (m, 1H), 1.40-1.60 (m, 3H).

10 **Example 141 :**

Preparation of 4-(7-Chloro-4-quinolinyl)-N-[3-(2-pyrimidinyloxy)cycloheptyl]-1-piperazinecarboxamide

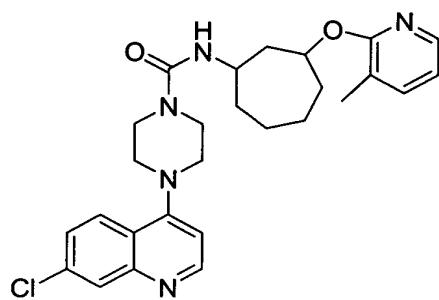


As described for example 138, 4-(7-chloro-4-quinolinyl)-N-(3-hydroxycycloheptyl)-1-piperazinecarboxamide, NaH, and 2-chloropyrimidine are reacted to give the product
15 after flash chromatography with CH₂Cl₂-MeOH. LC-MS: 481 (M⁺+1). ¹H NMR (CDCl₃) δ 8.75 (d, 1H), 8.50 (d, 2H), 8.05 (s, 1H), 7.95 (d, 1H), 7.45 (d, 1H), 6.92 (t, 1H), 6.85 (d, 1H), 5.55 (m, 1H), 5.15 (m, 1H), 4.10 (m, 1H), 3.70 (m, 4H), 3.23 (m, 4H), 1.80-2.25(m, 7H), 1.50-1.70 (m, 3H).

20

Example 142 :

Preparation of 4-(7-Chloro-4-quinolinyl)-N-[3-[(3-methyl-2-pyridinyl)oxy]cycloheptyl]-1-piperazinecarboxamide

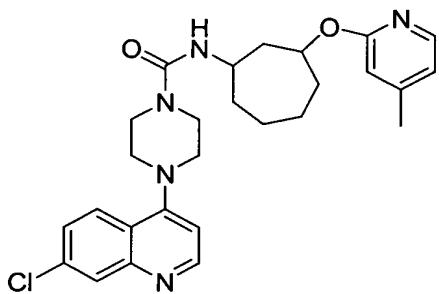


As described for example 138, 4-(7-chloro-4-quinolinyl)-N-(3-hydroxycycloheptyl)-1-piperazinecarboxamide, NaH, and 2-bromo-3-picoline are reacted to give the product.

5 LC-MS: 494 (M^++1). ^1H NMR (CDCl_3) δ 8.75 (d, 1H), 8.05 (s, 1H), 7.98 (d, 1H), 7.85 (m, 1H), 7.45 (d, 1H), 7.40 (m, 1H), 6.85 (d, 1H), 6.78 (m, 1H), 6.10 (d, 1H), 5.20 (m, 1H), 4.10 (m, 1H), 3.76 (m, 4H), 3.25 (m, 4H), 2.26 (m, 1H), 2.18 (s, 3H), 2.12 (m, 2H), 1.45-1.90 (m, 7H).

10 **Example 143 :**

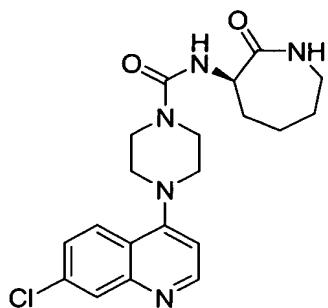
Preparation of 4-(7-Chloro-4-quinolinyl)-N-[3-[(4-methyl-2pyridinyl)oxy]cycloheptyl]-1-piperazinecarboxamide



15 As described for example 138, 4-(7-chloro-4-quinolinyl)-N-(3-hydroxycycloheptyl)-1-piperazinecarboxamide, NaH, and 2-bromo-4-picoline are reacted to give the product.
LC-MS 494 (M^++1). ^1H NMR (CDCl_3) δ 8.75 (d, 1H), 8.05 (s, 1H), 7.98 (d, 1H), 7.85 (d, 1H), 7.45 (d, 1H), 6.85 (d, 1H), 6.68 (d, 1H), 6.56 (s, 1H), 6.20 (d, 1H), 5.16 (m, 1H), 4.10 (m, 1H), 3.78 (m, 4H), 3.25 (m, 4H), 2.25 (s, 3H), 2.10 (m, 3H), 1.85 (m, 2H),
20 1.30-1.73 (m, 5H).

Example 144 :

Preparation of 4-(7-Chloro-4-quinolinyl)-N-[(3*R*)-hexahydro-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamid

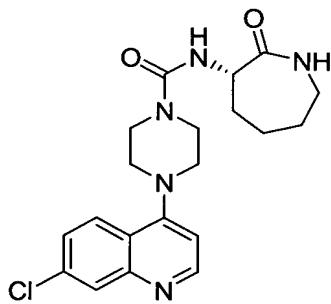


As described for example 78, (3*S*)-amino-azepin-2-one, *p*-nitrophenyl chloroformate,

5 diisopropylethyl amine, and 7-chloro-4-piperazinylquinoline are reacted to afford the product. LC-MS: 402 (M^++1). ^1H NMR (CDCl_3) δ 8.76 (d, 1H), 8.05 (s, 1H), 7.96 (d, 1H), 7.45 (d, 1H), 6.85 (d, 1H), 6.15 (d, 1H), 6.00 (m, 1H), 4.50 (m, 1H), 3.70 (m, 4H), 3.30 (m, 2H), 3.20 (m, 4H), 2.16 (m, 1H), 2.05 (m, 1H), 1.85 (m, 2H), 1.50 (m, 2H).

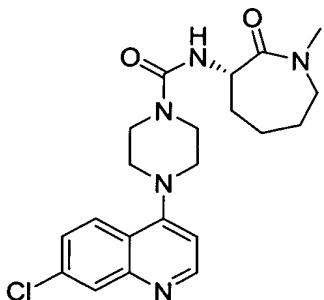
10 **Example 145 :**

Preparation of 4-(7-Chloro-4-quinolinyl)-N-[(3*S*)-hexahydro-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide



15 As described for example 78, (3*R*)-3-aminohexahydro-2*H*-azepin-2-one obtained by de-protection of [(3*R*)-hexahydro-2-oxo-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester with TFA, *p*-nitrophenyl chloroformate, diisopropylethyl amine and 7-chloro-4-piperazinylquinoline are reacted to afford the product. LC-MS 402 (M^++1). ^1H NMR (CDCl_3) δ 8.75 (d, 1H), 8.05 (s, 1H), 7.96 (d, 1H), 7.45 (d, 1H), 6.84 (d, 1H), 6.15 (d, 1H), 6.0 (m, 1H), 4.50 (m, 1H), 3.70 (m, 4H), 3.30 (m, 2H), 3.24 (m, 4H), 2.18 (m, 1H), 2.06 (m, 1H), 1.85 (m, 2H), 1.4-1.6 (m, 2H).

20

Example 146 :**Preparation of 4-(7-Chloro-4-quinolinyl)-N-[(3S)-hexahydro-1-methyl-2-oxo-1H-azepin-3-yl]-1-piperazinecarboxamide**

5 *Preparation of [(3R)-hexahydro-1-methyl-2-oxo-1H-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester:* To a stirred solution of [(3R)-hexahydro-2-oxo-1H-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester (500 mg) in THF (5 mL) at rt, KHDMS (5.26 mL, 1.0 M in THF) was added. After 30 min, CH₃I (0.26 mL) was introduced. The reaction was kept at rt overnight, and then quenched with brine. The reaction mixture
10 was extracted with EtOAc, washed with brine, and dried over Na₂SO₄. Concentration *in vacuo* afforded the crude product which was used directly for the next step reaction without further purification.

In a similar way, the following intermediates were prepared:

15

[(3S)-hexahydro-2-oxo-1-(phenylmethyl)-1H-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester

[(3S)-1-[(4-fluorophenyl)methyl]hexahydro-2-oxo-1H-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester

20 [(3S)-1-ethylhexahydro-2-oxo-1H-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester

[(3S)-1-(2-amino-2-oxoethyl)hexahydro-2-oxo-1H-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester

25 [(3S)-hexahydro-2-oxo-1-(3-pyridinylmethyl)-1H-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester

[(3S)-hexahydro-2-oxo-1-(2-propenyl)-1H-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester

[(3S)-1-(cyclopropylmethyl)hexahydro-2-oxo-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester

[(3S)-hexahydro-1-[(4-methoxyphenyl)methyl]-2-oxo-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester

5 (3S)-3-[[[(1,1-dimethylethoxy)carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-acetic acid, ethyl ester

(3S)-3-[[[(1,1-dimethylethoxy)carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-acetic acid, methyl ester

10 [(3S)-hexahydro-1-[3-(4-morpholinyl)propyl]-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester

[(3S)-1-[(2,6-dimethylphenyl)methyl]hexahydro-2-oxo-1*H*-azepin-3-yl]-carbamic acid-1,1-dimethylethyl ester

[(3S)-hexahydro-2-oxo-1-(2-pyridinylmethyl)-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester

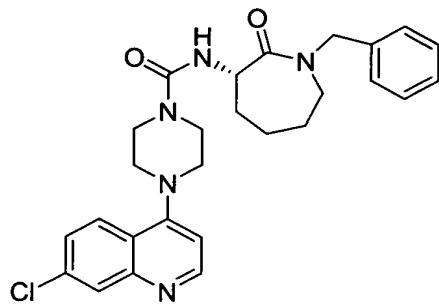
15 [(3S)-hexahydro-2-oxo-1-(4-pyridinylmethyl)-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester

*Preparation of 4-(7-chloro-4-quinoliny)-N-[(3S)-hexahydro-1-methyl-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide:* As described for example 78, (3*R*)-3-aminohexahydro-

20 1-methyl-2*H*-azepin-2-one obtained by de-protection of [(3*R*)-hexahydro-1-methyl-2-oxo-1*H*-azepin-3-yl]-carbamic acid-1,1-dimethylethyl ester with TFA, *p*-nitrophenyl chloroformate, diisopropylethyl amine and 7-chloro-4-piperazinylquinoline are reacted to afford the product. LC-MS 416 (M^++1). 1H NMR ($CDCl_3$) δ 8.78 (d, 1H), 8.05 (s, 1H), 7.95 (d, 1H), 7.46 (d, 1H), 6.85 (d, 1H), 6.25 (m, 1H), 4.60 (m, 1H), 3.68 (m, 4H), 2.30 (m, 4H), 1.80-2.10 (m, 5H), 1.45 (m, 3H).

Example 147 :

Preparation of 4-(7-Chloro-4-quinoliny)-N-[(3S)-hexahydro-2-oxo-1-(phenylmethyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide

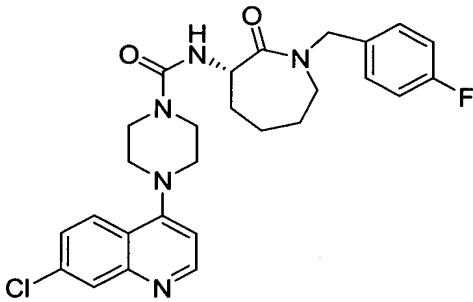


As described for example 78, (3*R*)-3-aminohexahydro-1-benzyl-2*H*-azepin-2-one obtained by de-protection of [(3*S*)-hexahydro-2-oxo-1-(phenylmethyl)-1*H*-azepin-3-yl]-

5 carbamic acid, 1,1-dimethylethyl ester with TFA, *p*-nitrophenyl chloroformate, diisopropylethyl amine and 7-chloro-4-piperazinylquinoline are reacted to afford the product. LC-MS 492 (M^++1). ¹H NMR ($CDCl_3$) δ 8.78 (d, 1H), 8.05 (s, 1H), 7.95 (d, 1H), 7.46 (d, 1H), 7.35 (m, 3H), 7.22 (m, 2H), 6.85 (d, 1H), 6.35 (m, 1H), 4.65 (m, 1H), 4.50 (m, 2H), 3.72 (m, 4H), 3.50 (m, 1H), 3.25 (m, 1H), 3.20 (m, 4H), 2.20 (m, 1H),
10 1.70-1.90 (m, 3H), 1.50 (m, 1H), 1.26 (m, 1H).

Example 148 :

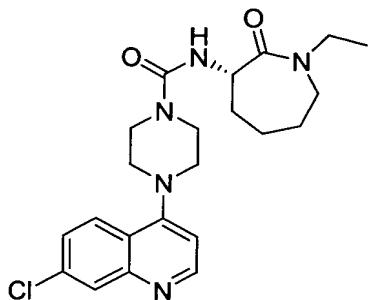
Preparation of 4-(7-Chloro-4-quinolinyl)-N-[(3*S*)-1-[(4-fluorophenyl)methyl]hexahydro-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide



15 As described for example 78, (3*R*)-3-amino-1-[(4-fluorophenyl)methyl]hexahydro-2*H*-azepin-2-one obtained by de-protection of [(3*S*)-1-[(4-fluorophenyl)methyl]hexahydro-2-oxo-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester with TFA, *p*-nitrophenyl chloroformate, diisopropylethyl amine, and 7-chloro-4-piperazinylquinoline are reacted to afford the product. LC-MS: 510 (M^++1). ¹H NMR ($CDCl_3$) δ 8.78 (d, 1H), 8.05 (s, 1H), 7.95 (d, 1H), 7.46 (d, 1H), 7.22 (m, 2H), 7.03 (t, 2H), 6.85 (d, 1H), 6.30 (m, 1H), 4.65 (m, 1H), 4.50 (m, 2H), 3.70 (m, 4H), 3.50 (m, 1H), 3.22 (m, 5H), 2.10 (m, 1H), 1.70-1.95 (m, 3H), 1.50 (m, 1H), 1.20 (m, 1H).

Example 149 :**Preparation of 4-(7-Chloro-4-quinoliny)-N-[(3S)-1-ethylhexahydro-2-oxo-1H-**

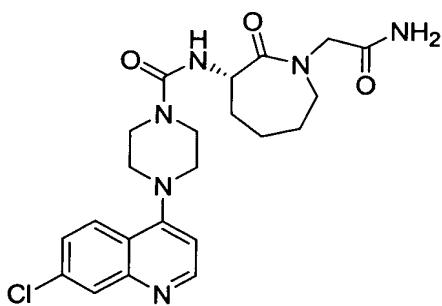
5 **azepin-3-yl]-1-piperazinecarboxamide**



As described for example 78, (3R)-3-amino-1-ethylhexahydro-2H-azepin-2-one obtained by de-protection of [(3S)-1-ethylhexahydro-2-oxo-1H-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester with TFA, *p*-nitrophenyl chloroformate, diisopropylethyl

10 amine, and 7-chloro-4-piperazinylquinoline are reacted to afford the product. LC-MS: 430 (M^++1). 1H NMR ($CDCl_3$) δ 8.78 (d, 1H), 8.05 (s, 1H), 7.95 (d, 1H), 7.46 (d, 1H), 6.85 (d, 1H), 6.30 (m, 1H), 4.58 (m, 1H), 3.70 (m, 4H), 3.50 (m, 2H), 3.42 (m, 1H), 3.22 (m, 5H), 2.10 (m, 1H), 1.85-1.98 (m, 3H), 1.46 (m, 2H), 1.17 (t, 3H).

15 **Example 150 :**

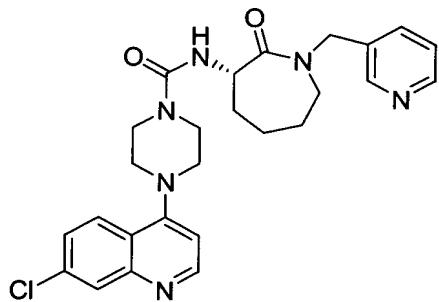
Preparation of (3S)-[[[4-(7-Chloro-4-quinoliny)-1-**piperazinyl]carbonyl]amino]hexahydro-2-oxo-1H-azepine-1-acetamide**

20 As described for example 78, (3R)-3-aminohexahydro-2-oxo-1H-azepine-1-acetamide obtained by de-protection of [(3S)-1-(2-amino-2-oxoethyl)hexahydro-2-oxo-1H-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester with TFA, *p*-nitrophenyl chloroformate, diisopropylethyl amine, and 7-chloro-4-piperazinylquinoline are reacted to afford the

product. LC-MS: 459 (M^++1). ^1H NMR (CDCl_3) δ 8.68 (d, 1H), 8.0 (s, 1H), 7.90 (d, 1H), 7.40 (d, 1H), 6.78 (d, 1H), 5.98 (m, 2H), 5.25 (m, 1H), 4.65 (m, 1H), 4.0 (m, 2H), 3.65 (m, 5H), 3.30 (m, 1H), 3.18 (m, 4H), 2.0 (m, 2H), 1.80 (m, 2H), 1.50 (m, 2H).

5 **Example 151 :**

Preparation of 4-(7-Chloro-4-quinolinyl)-N-[(3S)-hexahydro-2-oxo-1-(3-pyridinylmethyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide

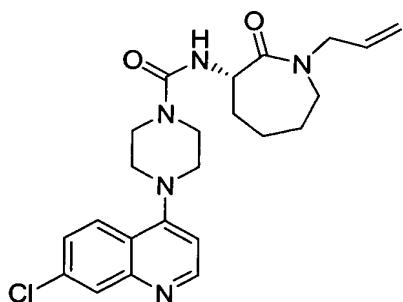


As described for example 78, (3*R*)-3-aminohexahydro-1-(3-pyridinylmethyl)-2*H*-

10 azepin-2-one obtained by de-protection of [(3*S*)-hexahydro-2-oxo-1-(3-pyridinylmethyl)-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester with TFA, *p*-nitrophenyl chloroformate, diisopropylethyl amine, and 7-chloro-4-piperazinylquinoline are reacted to afford the product. LC-MS: 493 (M^++1). ^1H NMR (CDCl_3) δ 8.70 (d, 1H), 8.50 (m, 2H), 8.05 (s, 1H), 7.92 (d, 1H), 7.60 (d, 1H), 7.40 (d, 1H), 7.22 (m, 1H), 6.80 (m, 1H), 6.20 (m, 1H), 4.65 (m, 1H), 4.56(m, 2H), 3.70 (m, 4H), 3.50 (m, 1H), 3.20 (m, 5H), 2.10 (m, 1H), 1.70-1.95 (m, 3H), 1.50 (m, 1H), 1.20 (m, 1H).

15 **Example 152 :**

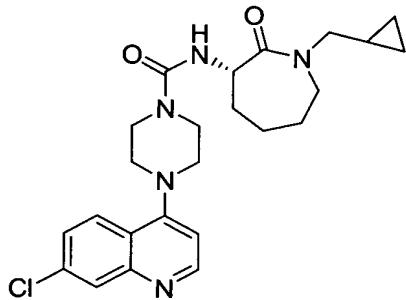
Preparation of 4-(7-Chloro-4-quinolinyl)-N-[(3S)-hexahydro-2-oxo-1-(2-propenyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide



As described for example 78, (3*R*)-3-aminohexahydro-1-(2-propenyl)-2*H*-azepin-2-one obtained by de-protection of [(3*S*)-hexahydro-2-oxo-1-(2-propenyl)-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester with TFA, *p*-nitrophenyl chloroformate, diisopropylethyl amine, and 7-chloro-4-piperazinylquinoline are reacted to afford the product. LC-MS: 442 (M^++1). 1H NMR ($CDCl_3$) δ 8.78 (d, 1H), 8.05 (s, 1H), 7.95 (d, 1H), 7.46 (d, 1H), 6.85 (d, 1H), 6.30 (m, 1H), 5.80 (m, 1H), 5.20 (m, 2H), 4.62 (m, 1H), 4.20 (m, 1H), 3.90 (m, 1H), 3.70 (m, 4H), 3.50 (m, 2H), 3.22 (m, 5H), 2.10 (m, 1H), 1.80-1.98 (m, 3H), 1.35-1.53 (m, 2H).

10 **Example 153 :**

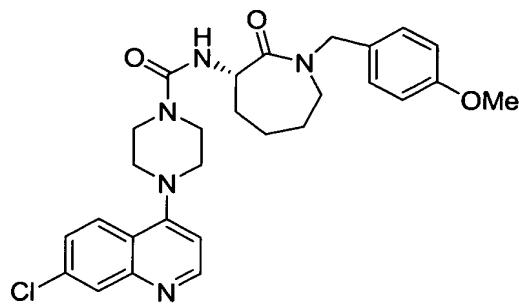
Preparation of 4-(7-Chloro-4-quinolinyl)-*N*-[(3*S*)-1-(cyclopropylmethyl)hexahydro-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide



As described for example 78, (3*R*)-3-amino-1-(cyclopropylmethyl)hexahydro-2*H*-azepin-2-one obtained by de-protection of [(3*S*)-1-(cyclopropylmethyl)hexahydro-2-oxo-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester with TFA, *p*-nitrophenyl chloroformate, diisopropylethyl amine, and 7-chloro-4-piperazinylquinoline are reacted to afford the product. LC-MS: 456 (M^++1). 1H NMR ($CDCl_3$) δ 8.78 (d, 1H), 8.05 (s, 1H), 7.95 (d, 1H), 7.46 (d, 1H), 6.85 (d, 1H), 6.30 (m, 1H), 4.62 (m, 1H), 3.70 (m, 4H), 3.61 (m, 1H), 3.46 (m, 2H), 3.26 (m, 5H), 2.10 (m, 1H), 1.80-2.00 (m, 3H), 1.56 (m, 2H), 0.96 (m, 1H).

Example 154 :

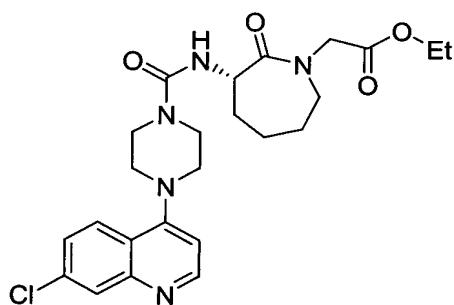
Preparation of 4-(7-Chloro-4-quinolinyl)-*N*-[(3*S*)-hexahydro-1-[(4-methoxyphenyl)methyl]-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide



As described for example 78, (3*R*)-3-aminohexahydro-1-[(4-methoxyphenyl)methyl]-2*H*-azepin-2-one obtained by de-protection of [(3*S*)-hexahydro-1-[(4-methoxyphenyl)methyl]-2-oxo-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester with TFA, *p*-nitrophenyl chloroformate, diisopropylethyl amine, and 7-chloro-4-piperazinylquinoline are reacted to afford the product. LC-MS: 552 (M^++1). ^1H NMR (CDCl_3) δ 8.70 (d, 1H), 8.05 (s, 1H), 7.90 (d, 1H), 7.40 (d, 1H), 7.15 (d, 2H), 6.80 (m, 2H), 6.30 (m, 1H), 4.60 (m, 1H), 4.50 (m, 2H), 3.76 (s, 3H), 3.65 (m, 4H), 3.40 (m, 1H), 3.20 (m, 5H), 2.10 (m, 1H), 1.60-1.90 (m, 3H), 1.50 (m, 1H), 1.20 (m, 1H).

Example 155 :

Preparation of (3*S*)-[[[4-(7-Chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-acetic acid, ethyl ester

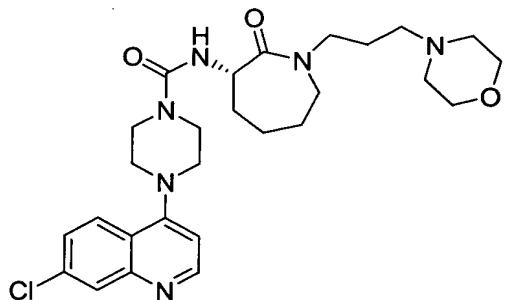


As described for example 78, (3*R*)-3-aminohexahydro-2-oxo-1*H*-azepine-1-acetic acid, ethyl ester obtained by de-protection of (3*S*)-3-[(1,1-dimethylethoxy)carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-acetic acid, ethyl ester with TFA, *p*-nitrophenyl chloroformate, diisopropylethyl amine, and 7-chloro-4-piperazinylquinoline are reacted to afford the product. LC-MS: 488 (M^++1). ^1H NMR (CDCl_3) δ 8.68 (d, 1H), 8.0 (s, 1H), 7.90 (d, 1H), 7.40 (d, 1H), 6.70 (d, 1H), 6.15 (d,

1H), 4.60 (m, 1H), 4.20 (m, 4H), 3.65 (m, 5H), 3.20 (m, 5H), 2.10 (m, 1H), 1.96 (m, 1H), 1.80 (m, 2H), 1.60 (m, 2H), 1.25 (t, 3H).

5 **Example 156 :**

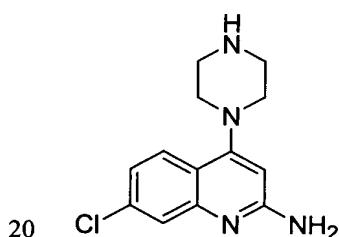
Preparation of 4-(7-Chloro-4-quinolinyl)-N-[(3S)-hexahydro-1-[3-(4-morpholinyl)propyl]-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide

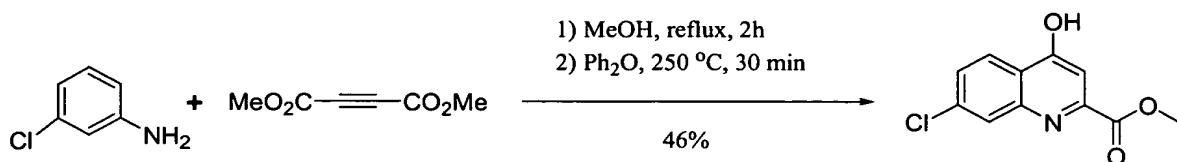


As described for example 78, (3*R*)-3-aminohexahydro-1-[3-(4-morpholinyl)propyl]-2*H*-azepin-2-one obtained by de-protection of [(3*S*)-hexahydro-1-[3-(4-morpholinyl)propyl]-2-oxo-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester with TFA, *p*-nitrophenyl chloroformate, diisopropylethyl amine, and 7-chloro-4-piperazinylquinoline are reacted to afford the product. LC-MS: 529 (M^++1). ^1H NMR (CDCl_3) δ 8.68 (d, 1H), 8.0 (s, 1H), 7.90 (d, 1H), 7.40 (d, 1H), 6.78 (d, 1H), 6.20 (d, 1H), 4.50 (m, 1H), 3.60 (m, 7H), 3.50 (m, 3H), 3.30 (m, 1H), 3.10 (m, 5H), 2.30-2.50 (m, 6H), 2.05 (m, 1H), 1.90 (m, 1H), 1.60- 1.80 (m, 4H), 1.25-1.40 (m, 2H).

Example 157 :

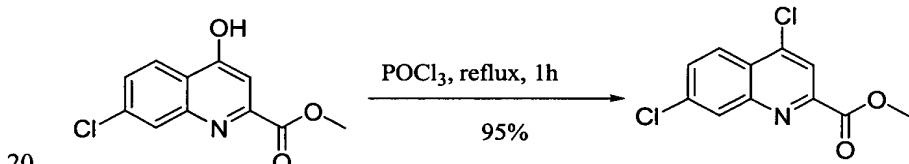
Preparation of 7-Chloro-4-(1-piperazinyl)-2-quinolinamine



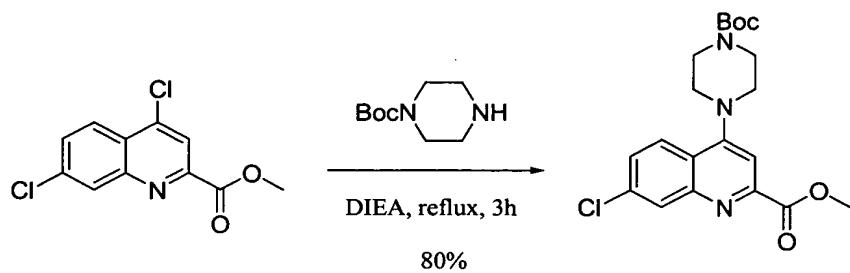


Preparation of 7-chloro-4-hydroxy-2-quinolinecarboxylic acid methyl ester [1. Juan C. Jaen, et al, J. Med. Chem., 1995, 38, 4439-4445. 2. WO 91/01973]:

To a stirred solution of 3-chloroaniline (40.8 g, 320 mmol) in MeOH (500 mL) at rt was added dimethyl acetylene dicarboxylate (50 g, 352 mmol). After addition, the reaction was kept at reflux for 2 h, then cooled to rt. After evaporating methanol *in vacuo*, the resulting crude adduct was dissolved in Ph₂O (160 mL), and was added dropwise to a hot solution of Ph₂O (800 mL) in a three-neck flask at 250 °C. The reaction mixture was kept at 250 °C for an additional 30 min, and was cooled to rt. The reaction mixture was diluted with hexane, and the product precipitated. The crude mixture was collected by filtration (washed with hexane and ether). This mixture (78 g) was re-dissolved in pyridine (468 mL) (Note: the other solvent for re-crystallization is acetic acid, and the ratio of crude product to acetic acid is 1: 6 based on weight), and kept at reflux for about 30 min. After cooling to rt, the desired product was collected by filtration (washed with ether). The solid was air dried overnight to give 35 g (46%) of 7-chloro-4-hydroxy-2-quinolinecarboxylic acid methyl ester. LC -MS: 237 (M⁺+1). ¹H NMR (DMSO-d₆): δ 3.98 (s, 3H), 6.6 (s, 1), 7.38 (d, 1H), 7.96 (s, 1H), 8.04 (d, 1), 12.1 (s, 1).

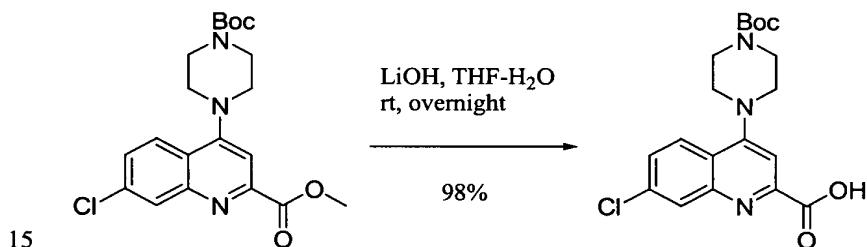


Preparation of 4,7-dichloro-2-quinolinecarboxylic acid methyl ester (D. J. Krogstad et al, J. Hetero. Chem., 1997, 34, 315-320): 7-Chloro-4-hydroxy-2-quinolinecarboxylic acid methyl ester (76 g) was added to POCl₃ (100 mL) at rt, and heated at 140 degrees C for 1 h, then cooled to rt. After the removal of excess POCl₃ *in vacuo*, cold 1N NaOH solution was added to pH 8, and precipitated solid was triturated, washed with water and dried under vacuum to afford 78 g (95.2%) of 4,7-dichloro-2-quinolinecarboxylic acid methyl ester as an off-white powder. LC-MS, 255 (M⁺+1). ¹H NMR (DMSO-d₆): δ 3.95 (s, 3H), 7.91 (d, 1H), 8.22 (s, 1H), 8.26 (d, 1H), 8.33 (s, 1H).



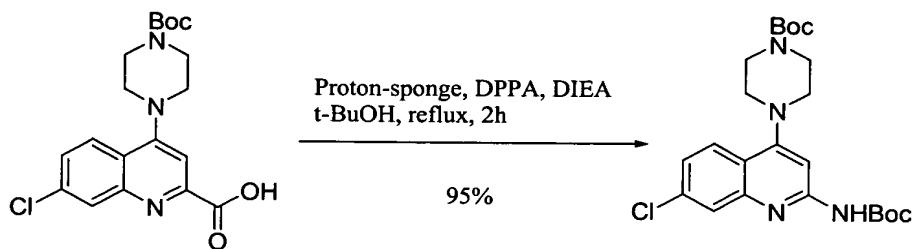
Preparation of 7-chloro-4-[4-[(1,1-dimethylethoxy)carbonyl]-1-piperazinyl]-2-quinolinecarboxylic acid methyl ester (D. J. Krogstad et al, *J. Hetero. Chem.*, 1997, 34,

5 315-320): The mixture of 4,7-dichloro-2-quinolinecarboxylic acid methyl ester (90.2 g, 379 mmol) and N-boc-piperazine (60.1 g, 387 mmol) in dry diisopropylethyl amine (300 mL) was kept at 140 degrees C for 3 h. After the removal of diisopropylethyl amine *in vacuo*, ethyl ether was added to precipitate the product. The crude solid was collected by filtration and was purified by re-crystallization from EtOAc (500 mL) to afford 123 g (80%) of 7-chloro-4-[4-[(1,1-dimethylethoxy)carbonyl]-1-piperazinyl]-2-quinolinecarboxylic acid methyl ester as an off-white solid. ^1H NMR (DMSO- d_6): δ 1.42 (s, 18H), 3.18 (m, 4H), 3.61 (m, 4H), 3.92 (s, 3H), 7.47 (s, 1 H), 7.63 (d, 1H), 8.04 (d, 1H), 8.1 (s, 1H).



15 *Preparation of 7-chloro-4-[4-[(1,1-dimethylethoxy)carbonyl]-1-piperazinyl]-2-quinolinecarboxylic acid:* To a stirred solution of 7-chloro-4-[4-[(1,1-dimethylethoxy)carbonyl]-1-piperazinyl]-2-quinolinecarboxylic acid methyl ester (100.4 g, 247 mmol) in THF (400 mL) was added LiOH• H_2O (15.6 g, 371 mmol) in water (400 mL) at rt. After stirring overnight, THF was removed *in vacuo*, and the resulting mixture was treated with 1N HCl to pH 4. The solid precipitated, and was collected by filtration, washed with water and cold ether. The solid was dried under vacuum overnight to afford 94.7 g (98%) of the product as a pale yellow solid. ^1H NMR (400 MHz, DMSO):

δ 1.39 (s, 9H), 3.28 (m, 4H), 3.59 (m, 4H), 7.51 (s, 1H), 7.63 (d, 1H), 8.07(d, 1H), 8.14 (s, 1H).

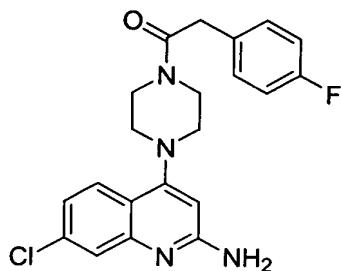


5 Preparation of 4-[7-chloro-2-[(1,1-dimethylethoxy)carbonyl]amino]-4-quinolinyl]- 1-piperazinecarboxylic acid-1,1-dimethylethyl ester[1] J. W. Gilman, et al, *Syn. Comm.*, 1993, 23, 335-341; 2) G. M. Coppola et al, *Bioorg. Med. Chem. Lett.*, 2000, 10, 1555-1558]: A mixture of 7-chloro-4-[4-[(1,1-dimethylethoxy)carbonyl]-1-piperazinyl]-2-quinolinecarboxylic acid (41 g, 105 mmol) proton-sponge (45 g, 210 mmol),
10 diphenylphosphoryl azide (DPPA) (37.7 g, 137 mmol), dry diisopropylethyl amine (DIEA) (17.7 g, 137 mmol) in t-BuOH (250 mL) was kept at reflux for 3 h, then cooled to rt. After the removal of the solvents *in vacuo*, the crude product was loaded onto silica-gel column, and washed with hexane-EtOAc (9:1) to afford 46.2 g (95%) of the product as an off-white solid. ^1H NMR (400 MHz, CDCl_3): δ 1.44 (s, 9H), 1.51 (s, 9H),
15 3.19 (m, 4H), 3.66 (m, 4H), 7.27 (d, 1H), 7.51 (s, 1H), 7.72 (s, 1H), 7.83 (d, 1H).

Preparation of 7-chloro-4-(1-piperazinyl)-2-quinolinamine: To a stirred sloution of 4-[7-chloro-2-[(1,1-dimethylethoxy)carbonyl]amino]-4-quinolinyl]-1-piperazinecarboxylic acid, 1,1-dimethylethyl ester (2 g, 4.33 mmol) in CH_2Cl_2 (30 mL) was added TFA (4 mL) at rt. The reaction mixture was kept at reflux for 4 h, then cooled to rt. After removal of the solvent and excess TFA, the mixture was diluted with CH_2Cl_2 , and treated with NaOH (10%) to pH 12. The mixture was extracted with CH_2Cl_2 , washed with brine, and dried over Na_2SO_4 . Concentration in vacuo afforded the product as a free base (1.13 g, 98%). LC-MS: 262 (M^++1). ^1H NMR (DMSO-d_6): δ 2.90 (s, 8H), 6.20
20 (s, 1H), 6.4 (s, 2H) 7.08 (d, 1H), 7.35 (s, 1H), 7.68 (d, 1H).
25 (s, 1H), 6.4 (s, 2H) 7.08 (d, 1H), 7.35 (s, 1H), 7.68 (d, 1H).

Example 158 :

Preparation of 7-Chloro-4-[4-[(4-fluorophenyl)acetyl]-1-piperazinyl]- 2-quinolinamine

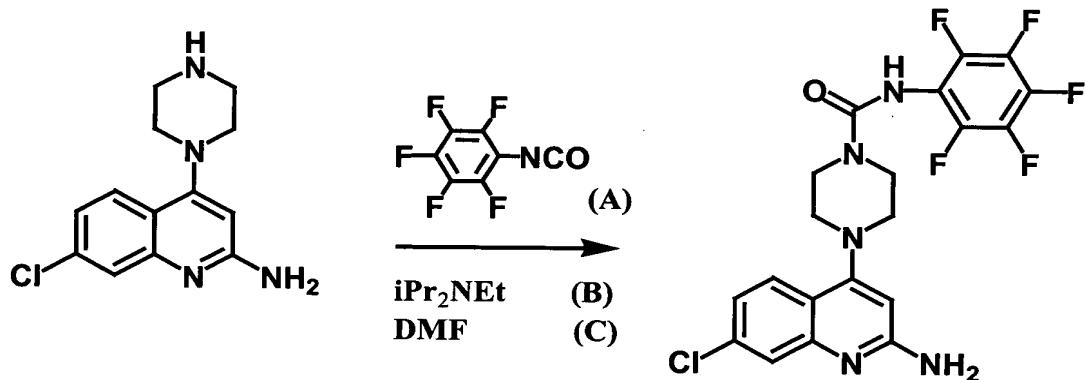


To a stirred solution of 4,7-dichloro-2-quinolinamine (50 mg, 0.19 mmol) in DMSO (10 mL) was added 4-fluorobenzeneacetic acid (32 mg, 0.2 mmol) followed by HATU (94 mg, 0.25 mmol) and triethylamine (0.053 ml, 0.38 mmol) at rt. After stirring overnight, 5 the reaction mixture was poured into ice-water (10 mL), extracted with EtOAc, washed with brine, and dried over Na_2SO_4 . Concentration in vacuo followed by preemptive TLC afforded the product. LC-MS: 398 (M^++1). ^1H NMR (DMSO- d_6): δ 2.9-3.05 (m, 4H), 3.68-3.85 (m, 6H), 6.22 (s, 1H), 6.45 (s, 2H), 7.05-7.2 (m, 3H), 7.22-7.33 (m, 2H), 7.35-7.4 (m, 1H), 7.7-7.8 (d, 1H).

10

Example 159 :

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-(2,3,4,5,6-pentafluorophenyl)-1-piperazinecarboxamide



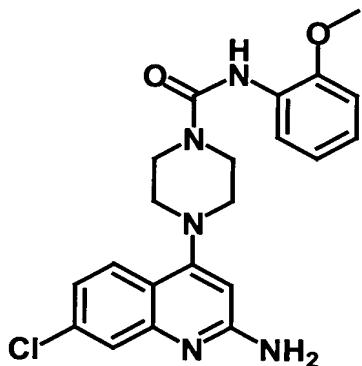
15

To a mixture of pentafluorophenyl isocyanate (0.03mL, 0.23mmol) and iPr_2NEt (0.05mL, 0.29 mmol) in DMF (1 mL), a hot solution of 7-chloro-4-(1-piperazinyl)-2-quinolinamine (50 mg, 0.19 mmol) in DMF (2 mL) was added. After stirring at rt overnight, the mixture was poured into ice-water. The solid was collected by filtration 20 and re-dissolved in CH_2Cl_2 -MeOH, then purified by preparative TLC (CH_2Cl_2 -MeOH, 9:1) affording the product as a light yellow solid. LC-MS: 472 (M^++1). ^1H NMR(DMSO-

d_6) δ 3.02 (br.s, 4H), 3.70 (br.s, 4H), 6.28 (s, 1H), 6.50 (s, 2H), 7.10 (d, 1H), 7.39 (s, 1H), 7.78 (d, 1H), 8.80 (s, 1H).

Example 160 :

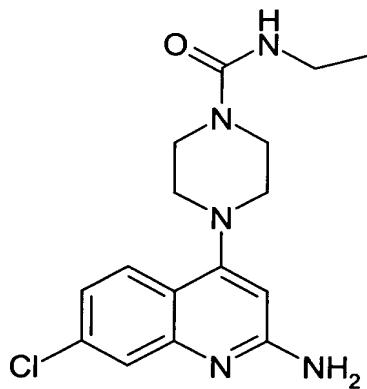
5 Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-(2-methoxyphenyl)-1-piperazinecarboxamide



As described for example 159, 7-chloro-4-(1-piperazinyl)-2-quinolinamine, 1-isocyanato-2-methoxy-benzene, and diisopropylethyl amine, are reacted to give the title product. LC-MS: 411(M^++1). ^1H NMR (CDCl_3) δ 3.15-3.25 (m, 4H), 3.7-3.85 (m, 4H), 3.9 (s, 3H), 4.6-4.8 (s, 2H), 6.2 (s, 1H), 6.86-6.9 (m, 1H), 6.94-7.02 (m, 2H), 7.16-7.21 (m, 2H), 7.62-7.65 (d, 1H), 7.74-7.78 (d, 1H), 8.14-8.19 (m, 1H).

Example 161 :

15 Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-ethyl-1-piperazinecarboxamide

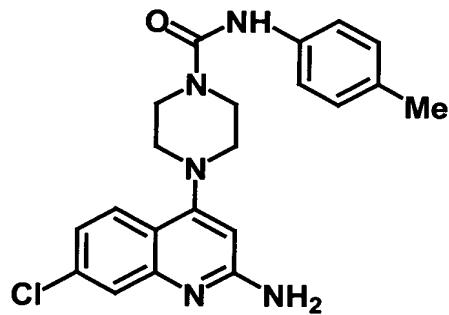


As described for example 159, 7-chloro-4-(1-piperazinyl)-2-quinolinamine, ethyl-isocyanate, and diisopropylethyl amine, are reacted to give the product.

LC-MS: 333 (M^++1). 1H NMR (DMSO-d₆): δ 1.0 (t, 3H), 2.8-3.0 (m, 4H), 3.0-3.14 (q, 2H), 3.45-3.6 (m, 4H), 6.25 (s, 1H), 6.4-6.55 (s, 2H), 6.55-6.65 (m, 1H), 7.1-7.2 (m, 1H), 7.35-7.4 (m, 1H), 7.7-7.8 (d, 1H).

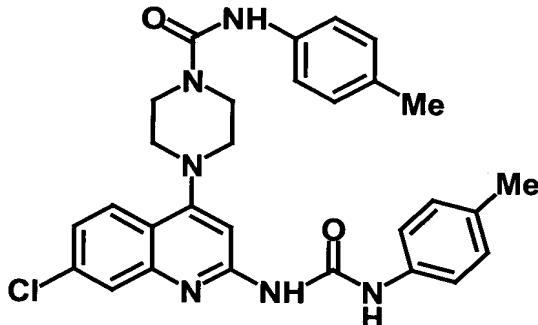
5 **Example 162:**

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-(4-methylphenyl)- 1-piperazinecarboxamide



As described for example 159, 7-chloro-4-(1-piperazinyl)-2-quinolinamine, 4-methylphenyl isocyanate, and diisopropylethyl amine, are reacted to give the product as a white solid. LC-MS: 395 (M^++1). 1H NMR (DMSO-d₆) δ 2.25 (s, 3H), 3.0-3.1(m, 4H), 3.64-3.75 (m, 4H), 6.3 (s, 1H), 6.5 (s, 2H), 7.0-7.1 (d, 2H), 7.1-7.2 (m, 1H), 7.3-7.45 (m, 3H), 7.75-7.85 (d, 1H), 8.5-8.6 (s, 1H).

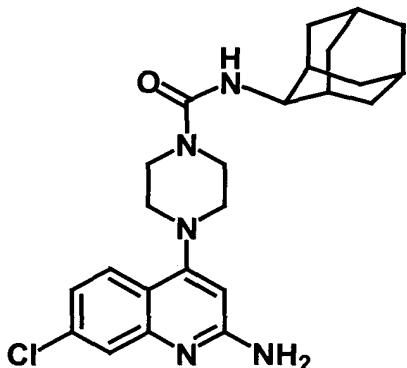
15 **4-[7-Chloro-2-[[[(4-methylphenyl)amino]carbonyl]amino]-4-quinolinyl]-N-(4-methylphenyl)-1-piperazinecarboxamide**



The title compound was obtained as product during the preparation of example 162. LC-MS: 529 ($M+1$). 1H NMR (DMSO-d₆) δ 2.18-2.35 (d, 6H), 3.1-3.3(m, 4H), 3.6-3.8(m, 4H), 6.8-6.9 (s, 1H), 7.0-7.1 (d, 2H), 7.1-7.2 (d, 2H), 7.3-7.48 (m, 3H), 7.55-7.64 (d, 2H), 7.9-8.0 (m, 2H), 8.5-8.6 (s, 1H), 9.8-9.9 (s, H), 11.62-11.69 (s, 1H).

Example 163 :

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-*N*-tricyclo[3.3.1.1^{3,7}]dec-2-yl-1-piperazinecarboxamide

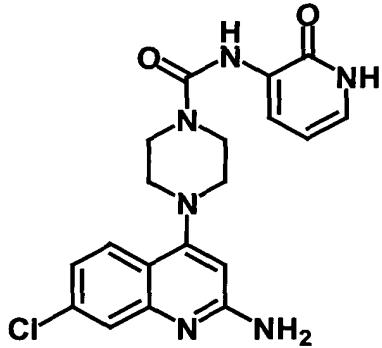


5 As described for example 159, 7-chloro-4-(1-piperazinyl)-2-quinolinamine, 1-admantyl isocyanate, and diisopropylethyl amine, are reacted to give the product. LC-MS: 440 (M+1). ¹H NMR (DMSO-d₆) δ 1.60, 1.95 and 2.11 (each s, 14H), 2.95 (br.s, 4H), 3.49 (m, 4H), 5.25 (m, 2H), 6.25 (s, 1H), 6.45 (br.s, 2H), 7.10 (dd, 1H), 7.38 (d, 1H), 7.74 (d, 1H).

10

Example 164 :

Preparation of 4-(2-amino-7-chloro-4-quinolinyl)-*N*-(1,2-dihydro-2-oxo-3-pyridinyl)-1-piperazinecarboxamide



15 *Preparation of 3-isocyanato-2(1H)-pyridinone:* To a solution of 3-amino-pyridone (220 mg, 2 mmol) in 1,4-dioxane (20 mL) was added diphosgene (0.363 mL) at 0 °C. After addition, the reaction mixture was stirred at rt for 2.5h, and the precipitate was formed. The solid was filtered, washed with Et₂O, dried *in vacuo* to afford the isocyanate (192 mg).

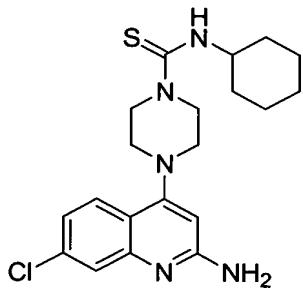
20

Preparation of 4-(2-amino-7-chloro-4-quinolinyl)-N-(1,2-dihydro-2-oxo-3-pyridinyl)-1-piperazinecarboxamide: As described example 159, 7-chloro-4-(1-piperazinyl)-2-quinolinamine, 3-isocyanato-2(1H)-pyridinone, and diisopropylethyl amine, are reacted to give the title product as a white solid. LC-MS: 399 (M^++1). 1H NMR (DMSO-d₆) δ 5 7.95 (m, 2H), 7.76(d, 1H), 7.40 (s, 1H), 7.12 (m, 1H), 7.0 (d, 1H), 6.45 (m, 2H), 6.25 (s, 1H), 6.22 (m, 1H), 3.75 (m, 4H), 3.10 (m, 4H).

Example 165 :

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-cyclohexyl-1-

10 **piperazinecarbothioamide**

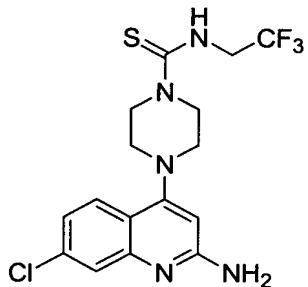


As described for example 159, 7-chloro-4-(1-piperazinyl)-2-quinolinamine, hexyl thioisocyanate, and diisopropylethyl amine, are reacted to give the product. LC-MS: 403 ($M+1$). 1H NMR (DMSO-d₆) δ 1.0-1.4 (m, 5H), 1.5-1.8 (m, 3H), 1.8-2.0 (m, 2H), 15 2.9-3.1 (m, 4H), 3.85-4.05 (m, 4H), 4.1-4.25 (m, 1H), 6.2-6.26 (s, 1H), 6.36-6.52 (s, 2H), 7.04-7.14 (m, 1 H), 7.32-7.44 (m, 2H), 7.7-7.78 (d, 1H).

Example 166 :

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-(2,2,2-trifluoroethyl)-1-

20 **piperazinecarbothioamide**

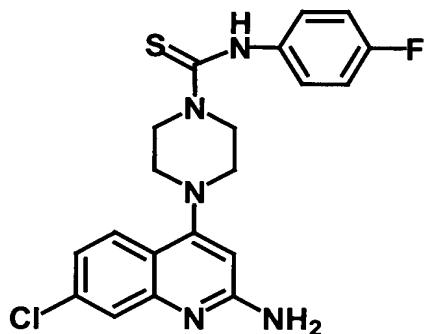


As described for example 159, 7-chloro-4-(1-piperazinyl)-2-quinolinamine, 1,1,1-trifluoro-2-isothiocyanato-ethane, and diisopropylethyl amine, are reacted to give the

product. LC-MS: 466 (M^++1). 1H NMR (DMSO-TFA-d₆) δ 1.0-1.3 (m, 1H), 3.0-3.14 (d, 1H), 3.18-3.38 (m, 6), 3.38-3.7 (m, 7H), 4.6-4.8 (m, 1H), 6.2-6.4 (s, 1H), 6.7-7.0 (m, 1H), 7.28-7.44 (m, 1H), 7.6-7.7 (s, 1H), 7.8-8.0 (m, 1H), 8.1-8.2 (m, 1H).

5 **Example 167 :**

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-(4-fluorophenyl)-1-piperazinecarbothioamide

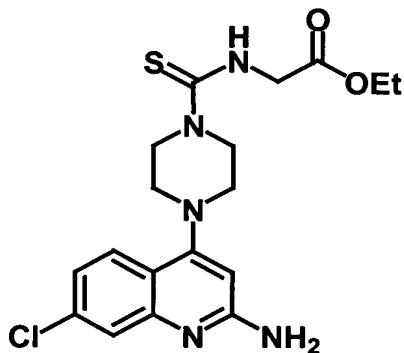


10 As described for example 159, 7-chloro-4-(1-piperazinyl)-2-quinolinamine, 4-fluorophenyl isothiocyanate, and diisopropylethyl amine, are reacted to give the product as a white solid. LC-MS: 416(M^++1). 1H NMR (DMSO-d₆) δ 3.10 (br.s, 4H), 4.15 (br.s, 4H), 6.20 (s, 1H), 6.45 (br.s, 2H), 7.10 (m, 3H), 7.30 (m, 2H), 7.40 (s, 1H), 7.80 (d, 1H), 9.42 (br.s, 1H).

15

Example 168 :

Preparation of [[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonothioyl]amino]-acetic acid-ethyl ester

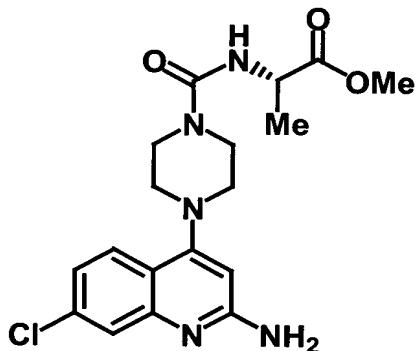


20 As described for example 159, 7-chloro-4-(1-piperazinyl)-2-quinolinamine, isothiocyanato-acetic acid-ethy ester, and diisopropylethyl amine, are reacted to give

the product as a white solid. LC-MS: 407 (M^++1). 1H NMR (DMSO-d₆): δ 1.16 (t, 3H), 3.05 (br.s, 4H), 4.01 (br.s, 4H), 4.06 (q, 2H), 4.20 (d, 2H), 6.24 (s, 1H), 6.47 (br.s, 2H), 7.08 (dd, 1H), 7.36 (d, 1H), 7.76 (d, 1H), 8.23 (dd, 1H).

5 **Example 169 :**

Preparation of (2S)-2-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-propanoic acid-methyl ester



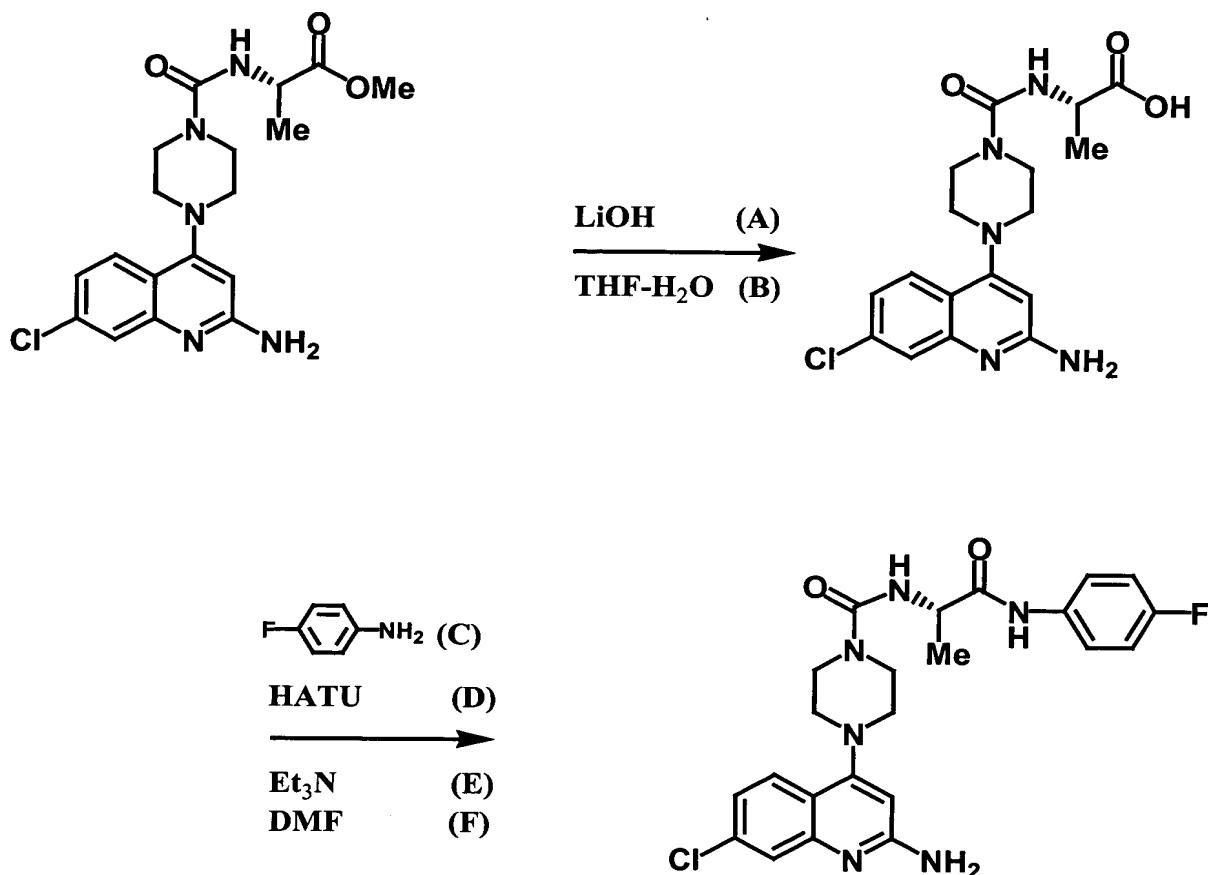
10

As described example 159, 7-chloro-4-(1-piperazinyl)-2-quinolinamine, 2-isocyanato-propanic acid-methyl ester, and diisopropylethyl amine, are reacted to give the product as a white solid. LC-MS: 392 (M^++1). 1H NMR (DMSO-d₆) δ 1.30 (d, 3H), 2.96 (br.s, 4H), 3.55 (m, 4H), 3.61 (s, 3H), 4.16 (m, 1H), 6.26 (s, 1H), 6.45 (br.s, 2H), 6.90 (d, 1H), 7.10 (dd, 1H), 7.38 (d, 1H), 7.75 (d, 1H).

15

Example 170 :

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(1S)-2-[(4-fluorophenyl)amino]-1-methyl-2-oxoethyl]-1-piperazinecarboxamide



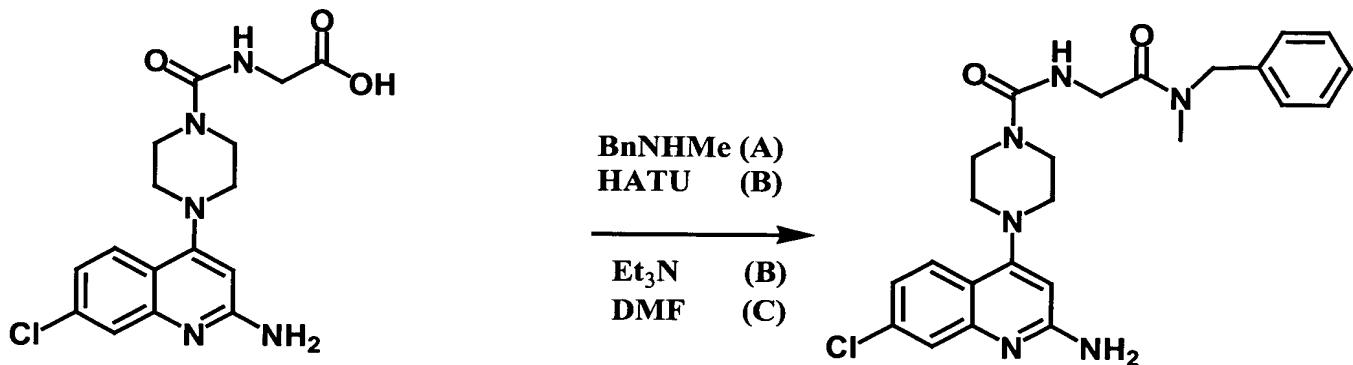
Preparation of (2S)-2-[[[4-(2-amino-7-chloro-4-quinoliny)-1-piperazinyl]carbonyl]amino]propanoic acid: A mixture of (2S)-2-[[[4-(2-amino-7-chloro-4-quinoliny)-1-piperazinyl]carbonyl]amino]-propanoic acid-methyl ester (~50 mg, 0.13 mmol), LiOH (100 mg) in THF-H₂O (5 mL, 5:1) was stirred at rt for 3h. The mixture was diluted with H₂O, and acidified with 1.0 N HCl to pH~5. The crude acid was obtained as a white solid by filtration.

Preparation of 4-(2-amino-7-chloro-4-quinoliny)-N-[(1S)-2-[(4-fluorophenyl)amino]-1-methyl-2-oxoethyl]-1-piperazinecarboxamide: To a solution of (2S)-2-[[[4-(2-amino-7-chloro-4-quinoliny)-1-piperazinyl]carbonyl]amino]-propanoic acid (~30mg), 4-fluoroalanine (0.03 mL, 0.32 mmol), Et₃N (0.14 mL, 0.35 mmol) in DMF (2mL) was added HATU(133 mg, 0.35 mmol) at rt . After stirring at rt overnight, the reaction mixture was poured into ice-water. The crude solid was obtained by filtration and re-dissolved in CH₂Cl₂-MeOH, dried over Na₂SO₄. Concentration *in vacuo* followed by preparative TLC(CH₂Cl₂-MeOH-Et₃N, 9:1:1%) afforded the title product as a light yellow powder. LC-MS: 471(M⁺+1). ¹H NMR (DMSO-d₆): δ 1.30 (d, 3H), 3.00 (br.s,

4H), 3.58 (m, 4H), 4.28 (m, 1H), 6.26 (s, 1H), 6.60 (br.s, 2H), 6.30 (d, 1H), 7.16 (m, 3H), 7.40 (d, 1H), 7.63 (dd, 2H), 7.75 (d, 1H).

Example 171 :

5 Preparation of 4-(2-Amino-7-chloro-4-quinoliny)-N-[2-[methyl(phenylmethyl)amino]-2-oxoethyl]-1-piperazinecarboxamide



Preparation of [[[4-(2-amino-7-chloro-4-quinoliny)-1-piperazinyl]carbonyl]amino]-acetic acid:

10 As described example 159, 7-chloro-4-(1-piperazinyl)-2-quinolinamine (1.5 g, 5.7 mmol), isocyanato-acetic acid-ethyl ester (0.7 mL, 6.3 mmol) and diisopropylethyl amine (1.5 mL, 8.6 mmol) are reacted to give [[[4-(2-amino-7-chloro-4-quinoliny)-1-piperazinyl]carbonyl]amino]-acetic acid-ethyl ester the product as a white solid (990 mg, 40%). A mixture of the ester (990 mg), LiOH.H₂O (1g) in THF-H₂O (5:1, 45 mL) was stirred at rt for 3h, then acidified with 1.0 N HC (aq.) to pH to 4~5. After the removal of THF *in vacuo*, crude product precipitated. 760 mg (76%) of the acid was obtained after drying under vacuum for 3 h. LC-MS: 363 (M⁺+1).

15

Preparation of 4-(2-amino-7-chloro-4-quinoliny)-N-[2-[methyl(phenylmethyl)amino]-2-

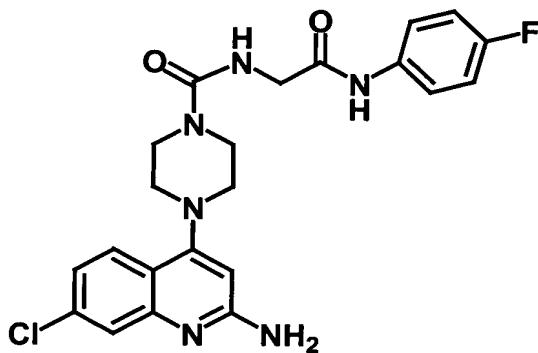
20 *oxoethyl]-1-piperazinecarboxamide:* To a mixture of [[[4-(2-amino-7-chloro-4-quinoliny)-1-piperazinyl]carbonyl]amino]-acetic acid (50 mg, 0.14 mmol), *N*-methylbenzylamine (0.022 mL, 0.17 mmol), iPr₂NEt (0.07 mL, 0.50 mmol) in DMF (3 mL) was added HATU (68 mg, 0.18 mmol) and stirred at rt overnight. The reaction mixture was poured into ice-water, and the solid was collected by filtration. The solid was re-dissolved in CH₂Cl₂ and purified by preparative TLC (CH₂Cl₂-MeOH-Hexane, 8:2:2) affording the product as a light yellow powder. LC-MS: 467(M⁺+1). ¹H NMR (CDCl₃): δ 2.92 and 3.00 (each s, 3H), 3.05~3.12 (m, 4H), 3.62~3.72 (m, 4H),

25

4.12~4.20 (m, 2H), 4.52 and 4.62 (each s, 2H), 5.08 (br.s, 2H), 5.94 (m, 1H), 6.15 (m, 1H), 7.14~7.24 (m, 3H), 7.26~7.40 (m, 3H), 7.58~7.62 (m, 1H), 7.68~7.72 (m, 1H).

Example 172 :

5 **Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[2-[(4-fluorophenyl)amino]-2-oxoethyl]-1-piperazinecarboxamide**

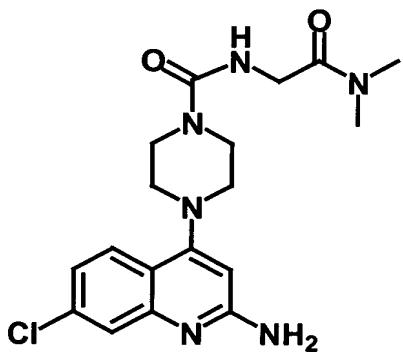


As described for example 171, [[[4-(2-amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-acetic acid, 4-fluorophenylaniline, iPr₂NE, HATU, are

10 reacted to afford the product. LC-MS: 456 (M⁺+1). ¹H NMR (CDCl₃): δ 3.05-3.3 (m, 4H), 3.6-3.8 (m, 4H), 4.0-4.2 (m, 2H), 4.65-4.9 (s, 2H), 6.1-6.2 (s, 1H), 7.0-7.1 (m, 2H), 7.2 (m, 1H), 7.3 (m, 2H), 7.6-7.7 (s, 1H), 7.7-7.8 (m, 1H), 8.6-8.7 (s, 1H).

Example 173 :

15 **Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[2-(dimethylamino)-2-oxoethyl]-1-piperazinecarboxamide**



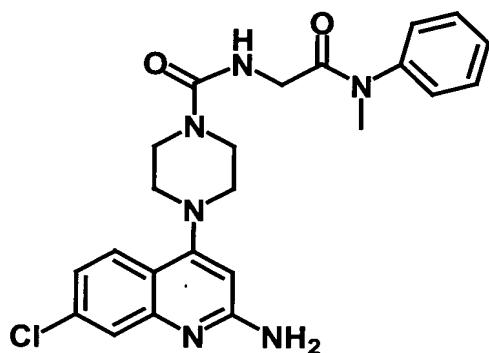
As described for example 171, [[[4-(2-amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-acetic acid, di-methylamine, iPr₂NEt, and HATU are

20 reacted to afford the product. LC-MS: 390 (M⁺+1). ¹H NMR (CDCl₃): δ 3.05 (s, 6H),

3.1-3.2 (m, 4H), 3.6-3.8 (m, 4H), 4.05-4.15 (d, 2H), 4.9-5.0 (s, H), 5.8 (m, 1H), 6.1-6.2 (s, 1H), 7.1-7.2 (m, 1H), 7.6 (m, 1H), 7.7-7.8 (m, 1H).

Example 174 :

5 **Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[2-(methylphenylamino)-2-oxoethyl]-1-piperazinecarboxamide**

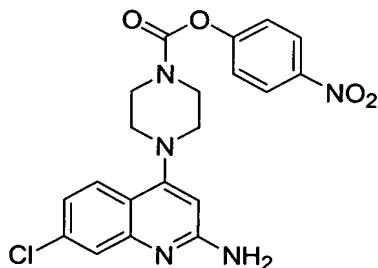


As described for example 171, [[[4-(2-amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-acetic acid, N-methyl-benzenamine, iPr₂NEt, and HATU

10 are reacted to afford the product. LC-MS: 452 (M⁺+1). ¹H NMR (CDCl₃): δ 3.0-3.2 (m, 4H), 3.3 (s, 3H), 3.6-3.76 (m, 4H), 3.76-3.9 (d, 2H), 4.8-5.0 (s, 2H), 5.6-5.8(m, 1H), 6.1-6.2 (s, 1H), 7.13-7.18 (m, 1H), 7.2-7.25 (m, 2H), 7.36-7.48 (m, 3H), 7.6 (d, 1H), 7.68-7.72 (d, 1H).

15 **Example 175 :**

4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinecarboxylic acid, 4-nitrophenyl ester

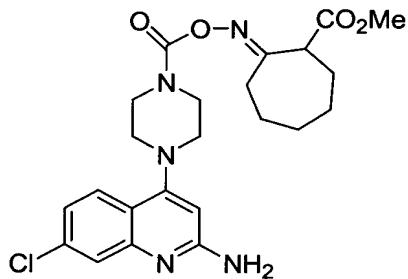


As described for example 78, 7-chloro-4-aminoquinoline, *p*-nitrophenyl chloroformate, 20 diisopropylethyl amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product. LC-MS: 427 (M⁺+1). ¹H NMR (DMSO-d₆) δ 3.0-3.2 (m, 4H), 3.6-3.9

(m, 4H), 6.2-6.3 (s, 1H), 6.4-6.6 (s, H), 7.05-7.2 (m, 1H), 7.35-7.4 (m, 1H), 7.4-7.5 (d, 2H), 7.7-7.8 (d, 1H), 8.2-8.35 (d, 2H).

Example 176 :

5 Preparation of 2-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]oxy]imino]-cycloheptanecarboxylic acid, methyl ester

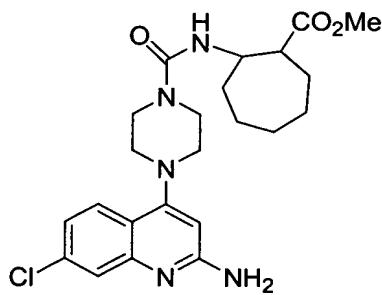


As described for 78, 2-methoxycarbonyl-cycloheptanone-oxime, 4-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product. LC-MS: 474 (M^++1). ^1H NMR (CDCl_3) δ 7.75 (d, 1H), 7.60 (s, 1H), 7.18 (d, 1H), 6.15 (s, 1H), 4.80 (s, 2H), 3.80 (m, 5H), 3.75 (s, 3H), 3.15 (m, 4H), 2.90 (m, 1H), 2.40 (m, 1H), 2.05 (m, 2H), 1.72 (m, 2H), 1.60 (m, 3H), 1.45 (m, 1H).

15

Example 177 :

Preparation of 2-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-cycloheptanecarboxylic acid, methyl ester

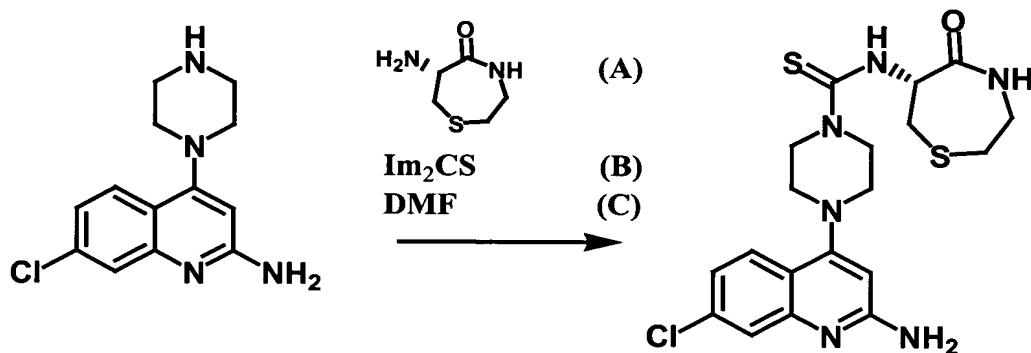


20 As described for 78, 2-methoxycarbonyl cycloheptyl amine, 4-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product. LC-MS: 460 (M^++1). ^1H NMR (CDCl_3) δ 7.72 (d, 1H), 7.60 (s, 1H), 7.18 (d, 1H), 6.15 (s, 1H), 5.50 (d, 1H), 4.70 (s, 2H), 4.15 (m, 1H), 3.70

(s, 3H), 3.60 (m, 4H), 3.10 (m, 4H), 2.95 (m, 1H), 2.00 (m, 1H), 1.85 (m, 3H), 1.65 (m, 5H), 1.45 (m, 1H).

Example 178 :

5 Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(6*R*)-hexahydro-5-oxo-1,4-thiazepin-6-yl]-1-piperazinecarbothioamide



10

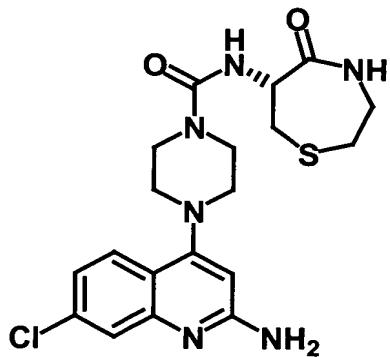
Preparation of (6*R*)-6-aminotetrahydro-1, 4-thiazepin-5(2*H*)-one (According to K. Soda, et al, *FEBS Letters*, 1984, 174, 76~79): To a mixture of L-cysteine methyl ester hydrochloride (5 g) in MeOH (30 mL) and Et₃N (7 mL) was added 2-chloroethylamine hydrochloride (5 g) in MeOH (10 mL) slowly at rt. After addition, the reaction mixture was kept at reflux for 16 h, then cooled to rt, and concentrated to give the crude product which was used for next step reaction without further purification.

Preparation of 4-(2-amino-7-chloro-4-quinolinyl)-N-[(6*R*)-hexahydro-5-oxo-1,4-thiazepin-6-yl]-1-piperazinecarbothioamide: To a solution of Im₂CS (41 mg, 0.29 mmol) in DMF (2 mL) was added slowly a suspension of (6*R*)-6-aminotetrahydro-1,4-thiazepin-5(2*H*)-one (34 mg, 0.23 mmol) in DMF (5 mL) at 0 °C. After 3 h at rt, the mixture was added to a solution of 7-chloro-4-(1-piperazinyl)-2-quinolinamine (50 mg, 0.19 mmol) in DMF (2 mL). After stirring at rt overnight, the reaction mixture was concentrated *in vacuo*, and the product precipitated. The title product was collected by filtration, and washed with chilly DMF and CH₂Cl₂. LC-MS: 451 (M⁺+1). ¹H NMR (DMSO-d₆): δ 2.50~2.56 (m, 1H), 2.62~2.74 (m, 2H), 2.80~2.86 (m, 1H), 3.40~3.52 (m,

6H), 4.02 (br.s, 4H), 5.30 (m, 1H), 6.25 (s, 1H), 7.40 (dd, 1H), 7.65~7.70 (m, 2H), 7.95 (d, 1H), 8.15 (m, 1H).

Example 179 :

5 **Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(6*R*)-hexahydro-5-oxo-1,4-thiazepin-6-yl]-1-piperazinecarboxamide**

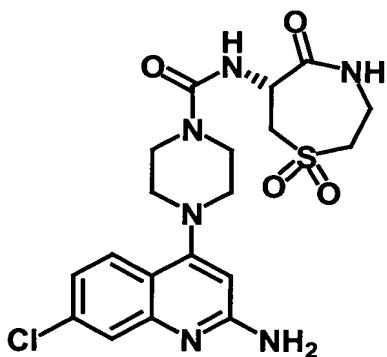


As described for example 78, (6*S*)-6-aminotetrahydro-1,4-thiazepin-5(2*H*)-one, *p*-nitrophenyl chloroformate, diisopropylethyl amine, and 7-chloro-4-(1-piperazinyl)-2-

10 quinolinamine are reacted to afford the product. LC-MS: 434 (M^++1). ^1H NMR (CDCl_3) δ 2.55-2.95 (m, 4), 3.0-3.25 (m, 4H), 3.6-3.9 (m, 6H), 4.8-5.0 (m, 3H), 6.1-6.2 (s, 1 H), 6.2-6.4 (m, 2H), 7.1-7.24 (m, 1H), 7.6 (s, 1H), 7.7-7.8 (m, 1H).

Example 180 :

15 **Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(6*R*)-hexahydro-1,1-dioxido-5-oxo-1,4-thiazepin-6-yl]-1-piperazinecarboxamide**



*Preparation of [(6*R*)-hexahydro-1,1-dioxido-5-oxo-1,4-thioazepin-6-yl]- carbamic acid,*

1,1-dimethylethyl ester: To a solution of (6*R*)-6-aminotetrahydro-1,4-thioazepin-5(2*H*)-

20 one (400 mg) in THF (50 mL) and brine (50 mL) was added $(\text{Boc})_2\text{O}$ (589 mg) at rt.

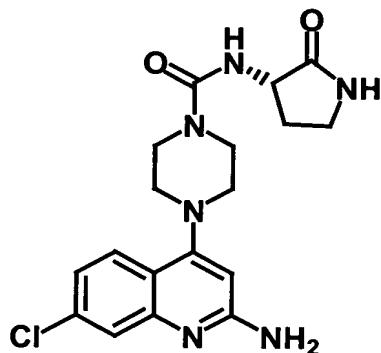
After 12 h, the reaction mixture was extracted with EtOAc, washed with water, and dried over Na₂SO₄. Concentration *in vacuo* afforded the crude [(6*S*)-hexahydro-5-oxo-1,4-thioazepin-6-yl]-carbamic acid, 1,1-dimethylethyl ester. This crude product was re-dissolved in MeOH, and treated with Oxone (3.71 g) for 24 h. After the removal of 5 most MeOH, the reaction mixture was diluted with water, and extracted with EtOAc, dried over Na₂SO₄. Concentration *in vacuo* gave the crude product which was used directly for next step reaction without further purification.

*Preparation of 4-(2-amino-7-chloro-4-quinolinyl)-N-[(6*R*)-hexahydro-1,1-dioxido-5-oxo-*

10 *1,4-thiazepin-6-yl]-1-piperazinecarboxamide:* As described for example 78, (6*R*)-6-aminotetrahydro-1,4-thioazepin-5(2*H*)-one, 1,1-dioxide obtained by de-protection of [(6*R*)-hexahydro-1,1-dioxido-5-oxo-1,4-thiazepin-6-yl]- carbamic acid, 1,1-dimethylethyl ester with TFA, *p*-nitrophenyl chloroformate, diisopropylethyl amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product. LC-MS: 15 466 (M⁺+1). ¹H NMR (DMSO-TFA-d₃) δ 1.0-1.3 (m, 1H), 3.0-3.14 (d, 1H), 3.18-3.38 (m, 6), 3.38-3.7 (m, 7H), 4.6-4.8 (m, 1H), 6.2-6.4 (s, 1H), 6.7-7.0 (m, 1H), 7.28-7.44 (m, 1H), 7.6-7.7 (s, 1H), 7.8-8.0 (m, 1H), 8.1-8.2 (m, 1H).

Example 181 :

20 **Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3*S*)-2-oxopyrrolidinyl]-1-piperazinecarboxamide**

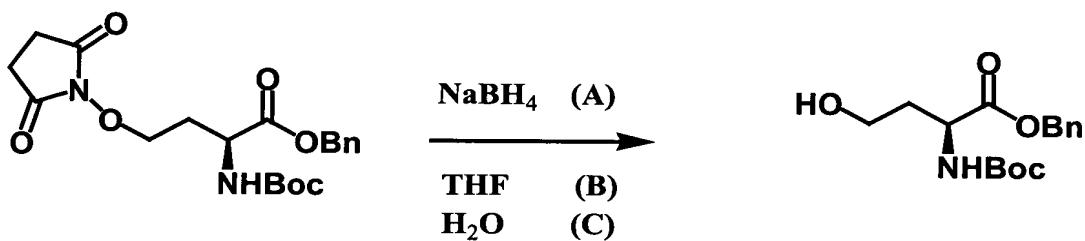
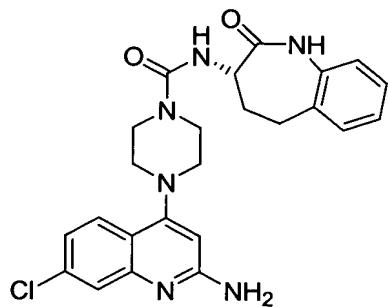


As described for example 78, (3*S*)-3-amino-2-pyrrolidinone, *p*-nitrophenyl chloroformate, diisopropylethyl amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine 25 are reacted to afford the product. LC-MS: 388 (M⁺+1). ¹H NMR (CDCl₃) δ 1.9-2.1 (m, 1), 2.8-2.9 (m, 1H), 3.0-3.2 (m, 4H), 3.35-3.5 (m, 2H), 3.6-3.8 (m, 4H), 4.2-4.4 (m, 1H),

4.7-5.0 (s, 2H), 5.3-5.4 (m, 1H), 5.7-5.85 (s, 1H), 6.1-6.2 (s, 1H), 7.1-7.2 (m, 1H), 7.6 (m, 1H), 7.7-7.8 (d, 1H).

5 Example 182 :

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-2,3,4,5-tetrahydro-2-oxo-1H-1-benzazepin-3-yl]-1-piperazinecarboxamide



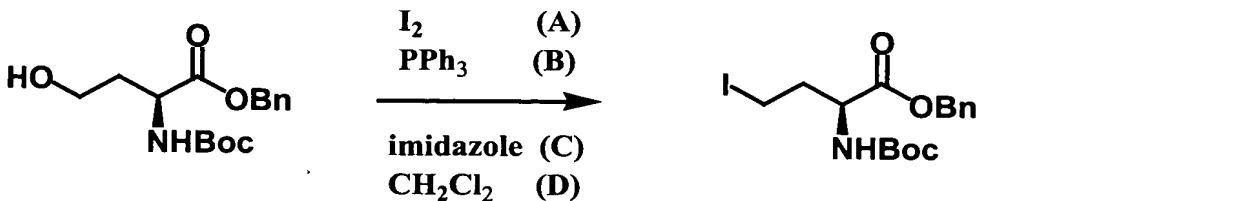
10

Preparation of (2S)-2-[[[(1,1-dimethylethoxy)carbonyl]amino]-4-hydroxy-butanoic acid phenylmethyl ester (Richard F. W. Jackson, Rebecca J. Moore, and Charles S.

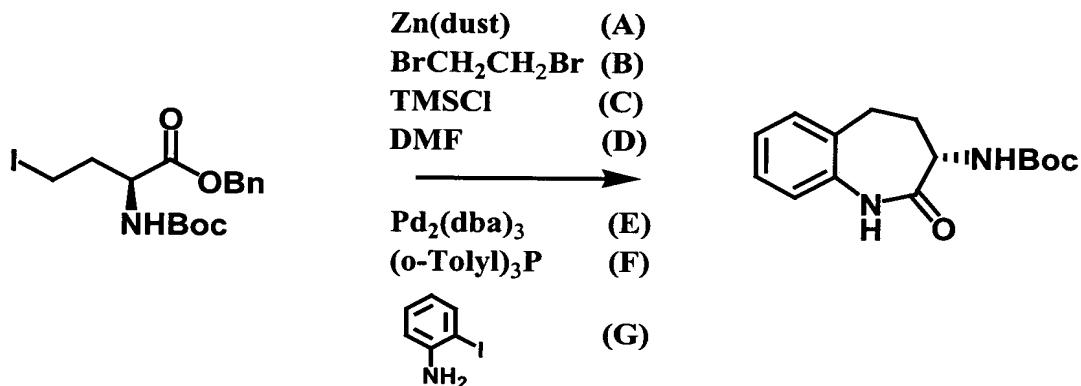
Dexter, Jason Elliott and Charles E Mowbray, *J. Org. Chem.*, **63**, 7875-7884(1998)):

To a mixture of NaBH₄ (1.4 g, 37 mmol) in THF (80 mL) and H₂O (20 mL) was added a solution of (2S)-2-[[[(1,1-dimethylethoxy)carbonyl]amino]-4-[(2,5-dioxo-1-pyrrolidinyl)oxy]-butanoic acid phenylmethyl ester (10 g, 24 mmol) in THF at 0 °C. After 15 min, the reaction mixture was quenched with NH₄Cl (sat.), and extracted with EtOAc, dried over Na₂SO₄. Concentration *in vacuo* followed by flash chromatography (hexane-EtOAc, 7:3 to 6:4) afforded the product as colorless oil. LC-MS: 310 (M⁺+1).

15

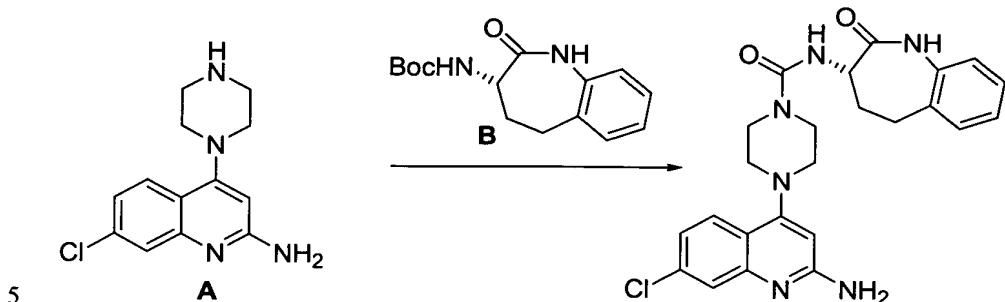


Preparation of (2S)-2-[(1,1-dimethylethoxy)carbonyl]amino]-4-iodo-butanoic acid phenylmethyl ester: To a mixture of PPh₃ (4.19 g, 16 mmol) and imidazole (1.09 g, 16 mmol) in anhydrous CH₂Cl₂ (40 mL) was added slowly iodine (4.07 g, 16 mmol) at rt. After 5 min, a solution of (2S)-2-[(1,1-dimethylethoxy)carbonyl]amino]-4-hydroxybutanoic acid phenylmethyl ester (4.12 g, 13.3 mmol) in CH₂Cl₂ (20 mL) was added at 0 °C. After 2h at rt, the reaction mixture was filtered through a short silica gel column. The filtrate was concentrated in vacuo, and the resulting residue was purified by flash chromatography (hexane-EtOAc, 95:5 to 9:1) to afford the product as light yellow oil (3.22g, 58%). ¹H NMR (CDCl₃) δ 1.43 (s, 9H), 2.20 (m, 1H), 2.40 (m, 1H), 3.13 (m, 2H), 4.35 (br.s, 1H), 5.10 (br.s, 1H), 5.20 (AB, 2H), 7.30~7.40 (m, 5H). LC-MS: 420 (M⁺+1).



15 *Preparation of [(3S)-2,3,4,5-tetrahydro-2-oxo-1H-1-benzazepin-3-yl]-carbamic acid-1,1-dimethylethyl ester:* To a nitrogen-purged flask with Zinc dust (2.5 g, 37.08 mmol) was added dry DMF (3 mL) and 1,2-dibromoethane (0.16 mL, 1.85 mmol). The reaction mixture was heated at 60 °C for 30min, then cooled to rt, and treated with trimethylsilyl chloride (0.05 mL, 0.39 mmol). After additional 30 min at rt, a solution of (2S)-2-[(1,1-dimethylethoxy)carbonyl]amino]-4-iodo-butanoic acid phenylmethyl ester (2.62 g, 6.25 mmol) in DMF (3 mL) was added. The reaction mixture was heated to 35°C for 30 min, then cooled to rt. To this reaction mixture, Pd₂(dba)₃ (110 mg, 0.12 mmol,), (o-Tolyl)₃P (144 mg, 0.47 mmol), and 2-iodoaniline (1.04 g, 4.74 mmol), were added sequentially. After 2 days at rt, the reaction mixture was quenched with brine, extracted with EtOAc, and dried over Na₂SO₄. Concentration *in vacuo* followed by flash chromatography (hexane-EtOAc, 9:1 to 8:2 to 7.5:2.5) afforded the product as a

light yellow solid (260 mg, 15%). LC-MS: 277 (M^++1). ^1H NMR (DMSO- d_6) δ 1.40 (s, 9H), 2.00 (m, 1H), 2.70 (m, 2H), 2.90 (m, 2H), 4.30 (m, 1H), 5.50 (m, 1H), 7.00 (d, 1H), 7.15 (m, 1H), 7.25 (m, 2H), 8.00 (br.s, 1H).



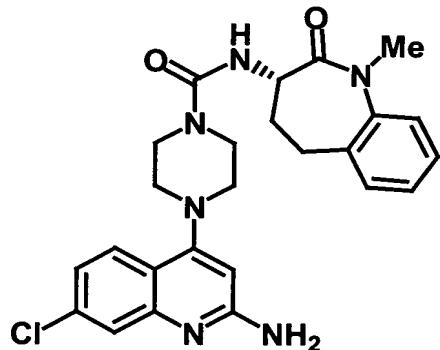
Preparation of 4-(2-amino-7-chloro-4-quinoliny)-N-[(3S)-2,3,4,5-tetrahydro-2H-1-benzazepin-3-yl]-1-piperazinecarboxamide: As described for example 78, (3R)-3-amino-1,3,4,5-tetrahydro-2H-1-benzazepin-2-one obtained by the de-protection of [(3S)-2,3,4,5-tetrahydro-2-oxo-1H-1-benzazepin-3-yl]-carbamic acid-1,1-dimethylethyl

10 ester with TFA, *p*-nitrophenyl chloroformate, diisopropylethyl amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product. ^1H NMR (CDCl₃) δ 2.02 (m, 1H), 2.67 (m, 1H), 2.81 (m, 1H), 2.98 (m, 1H), 3.12 (m, 4H), 3.64 (m, 4H), 4.52 (m, 1H), 4.96 (br. s, 2H), 5.69 (d, 1H), 6.17 (s, 1H), 6.98 (d, 1H), 7.17-7.3 (m, 5H), 7.58 (br. s, 1H), 7.6 (s, 1H), 7.72 (d, 1H).

15

Example 183 :

Preparation of 4-(2-Amino-7-chloro-4-quinoliny)-N-[(3S)-2,3,4,5-tetrahydro-1-methyl-2-oxo-1H-1-benzazepin-3-yl]-1-piperazinecarboxamide



20

Preparation of [(3R)-2,3,4,5-tetrahydro-1-methyl-2-oxo-1H-1-benzazepin-3-yl]-carbamic acid-1,1-dimethylethyl ester: As described for Example 146, [(3S)-2,3,4,5-tetrahydro-2-oxo-1H-1-benzazepin-3-yl]-carbamic acid-1,1-dimethylethyl ester, LiHMS, MeI are reacted to afford the product.

5

Preparation of 4-(2-amino-7-chloro-4-quinoliny)-N-[(3S)-2,3,4,5-tetrahydro-1-methyl-2-oxo-1H-1-benzazepin-3-yl]-1-piperazinecarboxamide: As described for example 78, (3R)-3-amino-1,3,4,5-tetrahydro-1-methyl-2H-1-benzazepin-2-one obtained by de-

protection of [(3R)-2,3,4,5-tetrahydro-1-methyl-2-oxo-1H-1-benzazepin-3-yl]-carbamic

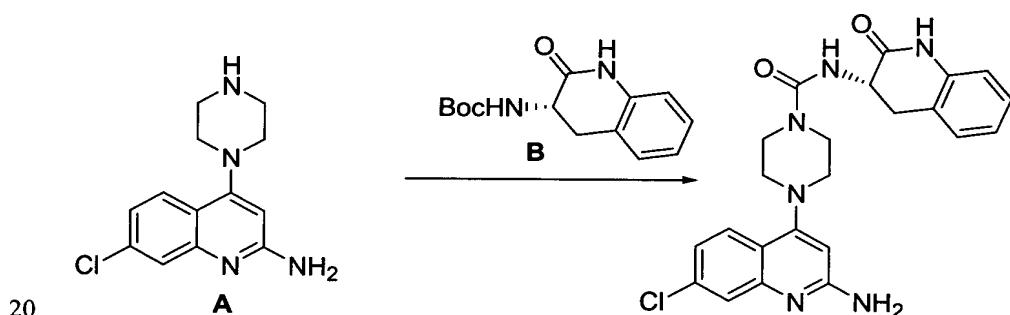
10 acid-1,1-dimethylethyl ester with TFA, *p*-nitrophenyl chloroformate, diisopropylethyl amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the

product. LC-MS: 479 (M^++1). ^1H NMR (CDCl_3): δ 1.90~2.00 (m, 1H), 2.60~2.74 (m, 2H), 2.80~2.90 (m, 1H), 3.10 (dd, 4H), 3.42 (s, 3H), 3.62 (dd, 4H), 4.40~4.48 (br.s, 2H), 5.72 (d, 1H), 6.13 (s, 1H), 7.14~7.26 (m, 4H), 7.30 (m, 1H), 7.61(d, 1H), 7.71 (d,

15 1H).

Example 184 :

Preparation of 4-(2-Amino-7-chloro-4-quinoliny)-N-[(3S)-1,2,3,4-tetrahydro-2-oxoquinoliny]-1-piperazinecarboxamide



20

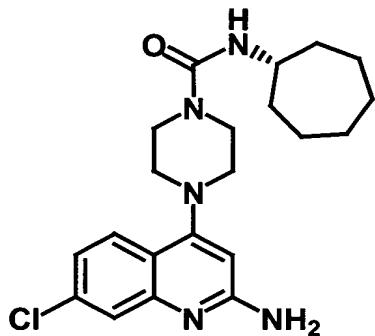
Preparation of [(3S)-1,2,3,4-tetrahydro-2-oxoquinoliny]-carbamic acid-1,1-dimethylethyl ester: The intermediate was prepared in a similar manner as for [(3S)-2,3,4,5-tetrahydro-2-oxo-1H-1-benzazepin-3-yl]-carbamic acid-1,1-dimethylethyl ester in Example 182. LC-MS: 263 (M^++1). ^1H NMR (DMSO-d_6): δ 1.40 (s, 9H), 2.80~3.00 (m, 2H), 4.10 (m, 1H), 6.80 (m, 1H) 6.90 (m, 1H), 6.95 (m, 1H), 10.20 (s, 1H).

Preparation of 4-(2-amino-7-chloro-4-quinolinyl)-N-[(3S)-1,2,3,4-tetrahydro-2-oxoquinolinyl]-1-piperazinecarboxamide: As described for example 78, (3R)-3-amino-3,4-dihydro-2(1H)-quinolinone obtained by de-protection of (3R)-1,2,3,4-tetrahydro-2-oxoquinolinyl]-carbamic acid, 1,1-dimethylethyl ester with TFA, *p*-nitrophenyl

5 chloroformate, diisopropylethyl amine, and 7-chloro-4-piperazinylquinoline are reacted to afford the product. ^1H NMR (CDCl_3) δ 2.82 (t, 1H), 3.17 (m, 4H), 3.58 (dd, 1H), 3.67 (m, 4H), 4.53 (m, 1H), 4.92 (br. s, 2H), 5.84 (br. s, 1H), 6.19 (s, 1H), 6.8 (d, 2H), 7.05 (t, 1H), 7.2 (m, 2H), 7.61 (s, 1H), 7.77 (d, 1H), 7.83 (br. s, 1H).

10 **Example 185 :**

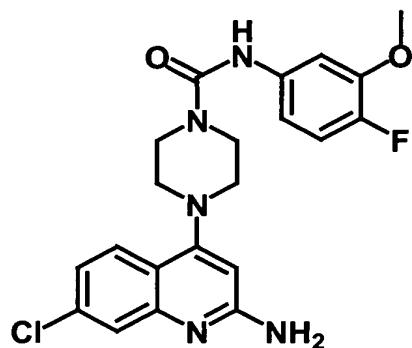
Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-cycloheptyl-1-piperazinecarboxamide



As described for example 78, cycloheptylamine, 4-nitrophenyl chloroformate,
15 diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a light yellow solid. LC-MS: 402 (M^++1). ^1H NMR (CDCl_3) δ 7.75 (d, 1H), 7.62 (s, 1H), 7.15 (d, 1H), 6.15 (s, 1H), 4.78 (s, 2H), 4.40 (d, 1H), 3.90 (m, 1H), 3.60 (m, 4H), 3.10(m, 4H), 1.95 (m, 2H) 1.40-1.65 (m, 10 H).

20 **Example 186 :**

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-(4-fluoro-3-methoxyphenyl)-1-piperazinecarboxamide

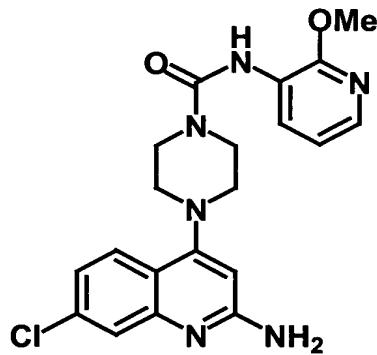


As described for example 78, 4-fluoro-3-methoxy-aniline, 4-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a light yellow solid. LC-MS: 430 (M^++1). 1H NMR (CD_3OD) δ

5 8.25 (m, 2H), 7.82 (m, 1H), 7.45 (s, 1H), 7.40 (d, 2H), 7.20 (d, 1H), 6.40 (s, 1H), 3.95 (s, 3H), 3.80 (m, 4H), 3.25 (m, 4H).

Example 187 :

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-(2-methoxy-3-pyridinyl)-1-piperazinecarboxamide

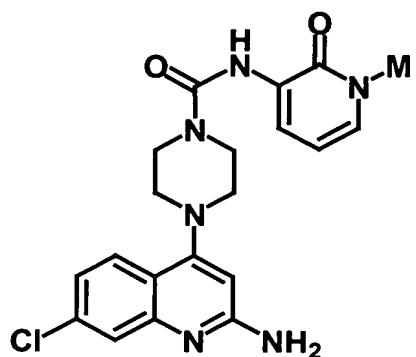


As described for example 78, 3-amino-2-methoxypyridine, 4-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a light yellow solid. LC-MS: 413 (M^++1). 1H NMR ($CDCl_3$) δ

15 8.40 (d, 1H), 7.80 (m, 1H), 7.75 (d, 1H), 7.64 (d, 1H), 7.20 (dd, 1H), 7.05 (s, 1H), 6.90 (m, 1H), 6.20 (s, 1H), 4.80 (s, 1H), 4.00 (s, 3H), 3.80 (m, 4H), 3.25 (m, 4H).

Example 188 :

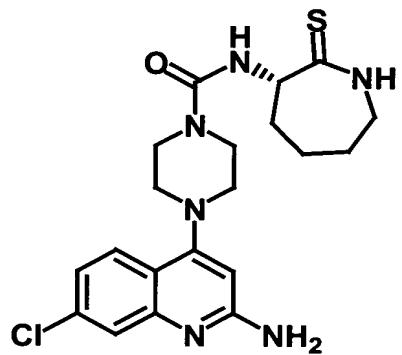
Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-(1,2-dihydro-1-methyl-2-oxo-3-pyridinyl)-1-piperazinecarboxamide



As described for example 78, 3-amino-1-methyl-2-pyridone, 4-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product. LC-MS: 413 (M^++1). ^1H NMR (CDCl_3) δ 8.15 (dd, 1H), 8.02 (s, 1H), 7.74 (d, 1H), 7.65 (s, 1H), 7.20 (dd, 1H), 6.95 (dd, 1H), 6.25 (m, 1H), 6.16 (s, 1H), 5.00 (s, 2H), 3.80 (m, 4H), 3.62 (s, 3H), 3.20 (m, 4H).

Example 189 :

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3*R*)-hexahydro-2-thioxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide



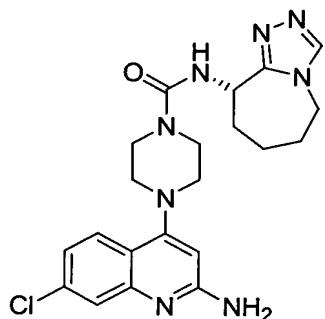
*Preparation of [(3*R*)-hexahydro-2-thioxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester: (3*S*)-Hexahydro-2-oxo-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester (1g, 4.24 mmol) and Lawesson's reagent (2.3 g) were mixed with toluene (10 mL) at reflux for 1h under nitrogen. After cooling to rt, the solvents was concentrated *in vacuo* and the resulting residue was purified by flash chromatography to yield the product.*

*Preparation of 4-(2-amino-7-chloro-4-quinolinyl)-N-[(3*R*)-hexahydro-2-thioxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide: As described for example 78, (3*R*)-3-*

aminohexahydro-2*H*-azepine-2-thione obtained by de-protection of [(3*R*)-hexahydro-2-thioxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester with TFA, 4-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a light yellow solid. LC-MS: 432 (M^++1). 1H NMR (CDCl₃) δ 1.2-1.65 (m, 2H), 1.8-2.2 (m, 4H), 3.05-3.3 (m, 4H), 3.35-3.5 (m, 1H), 3.56-3.9 (m, 5H), 4.6-4.8 (m, 1H), 4.8-5.1 (s, 2H), 6.1 (s, 1H), 6.8-7.0 (d, 1H), 7.14-7.22 (m, 1H), 7.6-7.65 (d, 1H), 7.7-7.78 (d, 1H), 8.56-8.74 (m, 1H).

Example 190 :

10 **Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(9S)-6,7,8,9-tetrahydro-5*H*-[1,2,4]triazolo[4,3-*a*]azepin-9-yl]-1-piperazinecarboxamide**



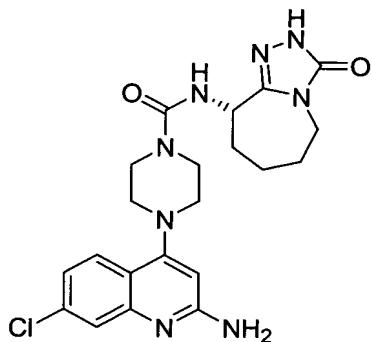
15 *Preparation of [(9S)-6,7,8,9-tetrahydro-5*H*-[1,2,4]triazolo[4,3-*a*]azepin-9-yl]- carbamic acid, 1,1-dimethylethyl ester:* To a mixture [(3*S*)-hexahydro-2-thioxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester and formic hydrazine (60 mg) in MeCN (6 mL) was added Hg(OAc)₂ (318 mg) at rt. After 2h, the reaction mixture was diluted with brine, extracted with EtOAc, and dried. Concentration in vacuo followed by flash chromatography gave the product.

20 *Preparation of 4-(2-amino-7-chloro-4-quinolinyl)-N-[(9S)-6,7,8,9-tetrahydro-5*H*-[1,2,4]triazolo[4,3-*a*]azepin-9-yl]-1-piperazinecarboxamide:* As described for example 78, (9*R*)-6,7,8,9-tetrahydro-5*H*-[1,2,4]triazolo[4,3-*a*]azepin-9-amine obtained by de-protection of [(9S)-6,7,8,9-tetrahydro-5*H*-[1,2,4]triazolo[4,3-*a*]azepin-9-yl]-carbamic acid, 1,1-dimethylethyl ester with TFA, 4-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product. LC-MS: 441 (M^++1). 1H NMR (CD₃OD) δ 1.20 (m, 2H), 1.50 (m, 1H), 1.70 (m, 1H), 1.80 (m, 1H), 2.00 (m, 3H), 3.10 (m, 4H), 3.70 (m, 4H), 3.90 (m,

1H), 4.35 (m, 1H), 5.05 (d, 1H), 6.30 (s, 1H), 7.10 (d, 1H), 7.40 (s, 1H), 7.80 (d, 1H), 8.35 (s, 1H).

Example 191 :

5 Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(9S)-2,5,6,7,8,9-hexahydro-3-oxo-3H-[1,2,4]triazolo[4,3- α]azepin-9-yl]-1-piperazinecarboxamide



Preparation of [(9S)-2,5,6,7,8,9-hexahydro-3-oxo-3H-[1,2,4]triazolo[4,3- α]azepin-9-yl]-

10 *carbamic acid, 1,1-dimethylethyl ester:* To a mixture of [(3S)-hexahydro-2-thioxo-1H-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester (300mg, 1.22 mmol) and methyl hydrazinocarboxylate (130 mg, 1.75 mmol) in MeCN (5 mL) was added Hg(OAc)₂ (510 mg), and the reaction mixture was stirred at rt for 2h. The solid was filtered off, and the filtrate was diluted with MeCN (10 mL), then K₂CO₃ was added. The reaction mixture was kept 80 °C for 30 min. After cooling to rt, the reaction mixture was diluted with AcOEt (250 mL), washed with water (100mL×2), and dried. Concentration *in vacuo* afforded the product.

15

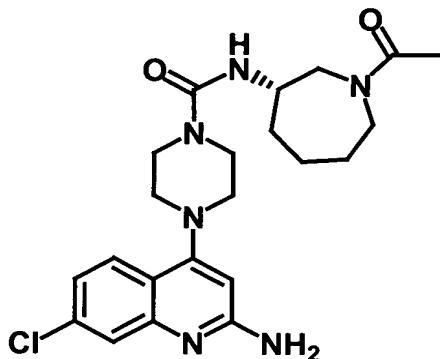
Preparation of 4-(2-amino-7-chloro-4-quinolinyl)-N-[(9S)-2,5,6,7,8,9-hexahydro-3-oxo-

20 *3H-[1,2,4]triazolo[4,3- α]azepin-9-yl]-1-piperazinecarboxamide:* As described for example 78, (9R)-9-amino-2,5,6,7,8,9-hexahydro-3H-[1,2,4]triazolo[4,3- α]azepin-3-one obtained by de-protection of [(9S)-2,5,6,7,8,9-hexahydro-3-oxo-3H-[1,2,4]triazolo[4,3- α]azepin-9-yl]- carbamic acid, 1,1-dimethylethyl ester with TFA, 4-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product. LC-MS: 457 (M⁺+1). ¹H NMR (CD₃OD) δ 1.75 (m,

3H), 2.0 (m, 3H), 3.20 (m, 1H), 3.40 (m, 5H), 3.75 (m, 4H), 4.10 (m, 1H), 6.37 (s, 1H), 7.30 (s, 1H), 7.40 (d, 1H), 7.60 (s, 1H), 7.95 (d, 1H).

Example 192 :

5 **Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)- N-[(3S)-1-acetylhexahydro-1*H*-azepin-3-yl]-1-piperazinecarboxamide**

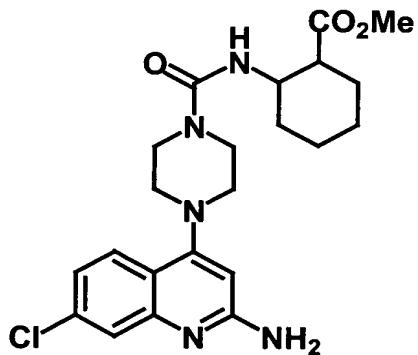


As described for example 78, (3*R*)-1-acetylhexahydro-1*H*-azepin-3-amine, 4-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a light yellow solid. LC-MS: 445 (M^++1). ^1H NMR (CDCl_3) δ 7.70 (d, 1H), 7.60 (s, 1H), 7.15 (d, 1H), 6.85 (d, 1H), 6.02 (s, 1H), 4.70 (s, 2H), 4.22 (d, 1H), 4.10 (m, 1H), 3.92 (m, 1H), 3.60 (m, 4H), 3.10 (m, 6H), 2.25 (m, 1H), 2.15(s, 3H), 1.60-1.90 (m, 3H), 1.30-1.40 (m, 2H).

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Example 193 :

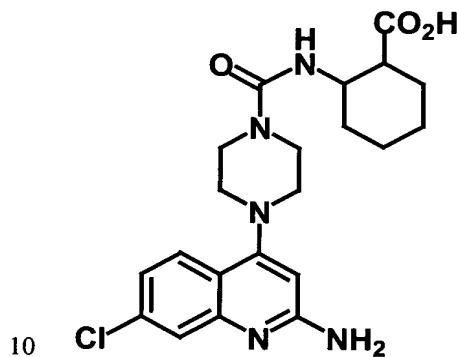
Preparation of 3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-cyclohexanecarboxylic acid, methyl ester



As described for example 78, 2-amino-cyclohexanecarboxylic acid-methyl ester, 4-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a light yellow solid. LC-MS: 446 (M^++1). 1H NMR (CD_3OD) δ 1.20-1.40 (m, 4H), 1.90 (m, 3H), 2.10 (m, 1H), 2.45 (m, 5 H), 3.30 (m, 4H), 3.70(m, 8H), 6.35 (s, 1H), 7.38 (d, 1H), 7.60 (s, 1H), 790 (d, 1H).

Example 194 :

Preparation of 3-[[[4-(2-Amino-7-chloro-4-quinoliny)-1-piperazinyl]carbonyl]amino]-cyclohexanecarboxylic acid

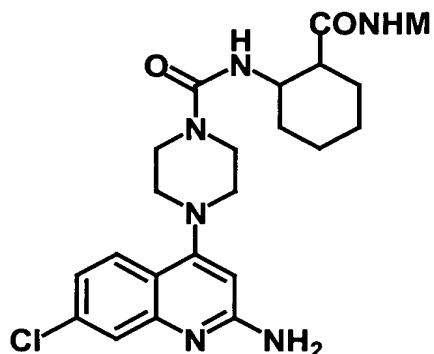


To a solution of 3-[[[4-(2-amino-7-chloro-4-quinoliny)-1-piperazinyl]carbonyl]amino]-cyclohexanecarboxylic acid, methyl ester (150 mg, 0.33 mmol) in THF (20 mL) and water (6 mL) was added LiOH. H_2O (28.0 mg). The reaction mixture was stirred at rt overnight, and then solvent was evaporated *in vacuo*. The resulting residue was acidified to pH4, the precipitate was collected by filtration and dried to afford the title product. LC-MS: 432 (M^++1). 1H NMR ($DMSO-d_6$) δ 1.10 (m, 2H), 1.25 (m, 2H), 1.75 (m, 3H), 1.95 (m, 1H), 2.25 (m, 1H), 3.10 (m, 4H), 3.50 (m, 5H), 6.25 (s, 1H), 6.40 (d, 1H), 7.25 (m, 1H), 7.50 (s, 1H), 780 (d, 1H).

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Example 195 :

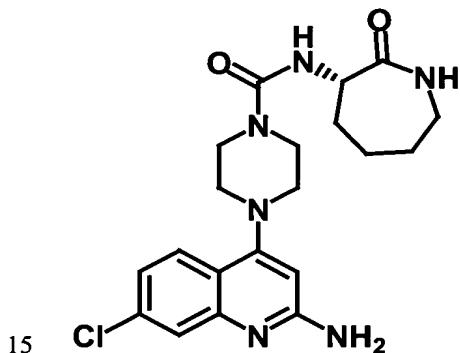
Preparation of 4-(2-Amino-7-chloro-4-quinoliny)-N-[3-[(methylamino)carbonyl]cyclohexyl]-1-piperazinecarboxamide



The mixture of 3-[[[4-(2-amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]cyclohexanecarboxylic acid (30 mg, 0.069 mmol), HATU (100 mg), methylamine (1.0M
5 in THF, 3 mL) and diisopropylethylamine (0.5mL) in DMF (3 mL) was stirred at rt for
3h. The reaction mixture was diluted with ethyl acetate (100 mL), washed with sat.
NaHCO₃ solution and water, and dried over Na₂SO₄. Concentration *in vacuo* followed
by flash chromatography afforded the title product. LC-MS: 445 (M⁺+1). ¹H NMR
(CD₃OD) δ 1.10-1.35 (m, 4H), 1.65-1.90 (m, 4H), 2.20 (m, 1H), 2.65 (s, 3H), 3.08 (m,
10 4H), 3.60 (m, 5H), 5.80 (d, 1H), 6.15 (s, 1H), 7.10 (d, 2H), 7.45 (s, 1H), 7.65 (d, 1H).

Example 196 :

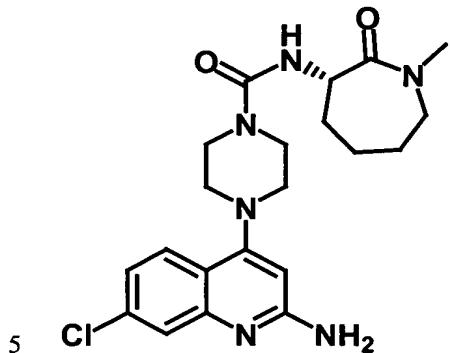
Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-hexahydro-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide



As described for example 78, (s)-3-amino-azepan-2-one, 4-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a light yellow solid. LC-MS: 417 (M⁺+1). ¹H NMR (CD₃OD) δ 7.80 (d, 1H), 7.45 (s, 1H), 7.15 (d, 1H), 6.35 (s, 1H), 4.45 (d, 1H), 3.68 (m, 4H), 3.25
20 (m, 2H), 3.10 (m, 4H), 2.00 (m, 2H), 1.80 (m, 2H), 1.60 (m, 1H), 1.38 (m, 1H).

Example 197 :

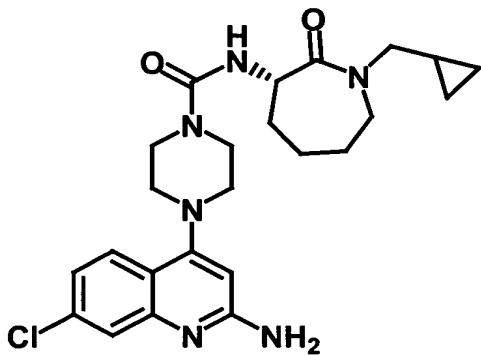
Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-hexahydro-1-methyl-2-oxo-1H-azepin-3-yl]-1-piperazin carboxamide



As described for example 78, (3*R*)-3-aminohexahydro-1-methyl-2*H*-azepin-2-one obtained by de-protection of [(3*R*)-hexahydro-1-methyl-2-oxo-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester with TFA, 4-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a light yellow solid. LC-MS: 431 (M^++1). ^1H NMR (CDCl_3) δ 7.70 (d, 1H), 7.60 (s, 1H), 7.15 (d, 1H), 6.25 (d, 1H), 6.15 (s, 1H), 4.95 (s, 2H), 4.60 (m, 1H), 3.68 (m, 5H), 3.20 (m, 1H), 3.10 (m, 4H), 3.05 (s, 3H), 2.10 (m, 1H), 1.95 (m, 1H), 1.85 (m, 2H), 1.46 (m, 2H).

15 Example 198 :

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-1-(cyclopropylmethyl)hexahydro-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide

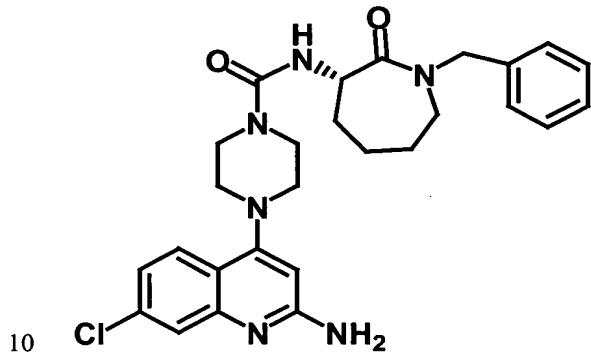


As described for example 78, (3*R*)-3-amino-1-(cyclopropylmethyl)hexahydro-2*H*-azepin-2-one by de-protection of [(3*S*)-1-(cyclopropylmethyl)hexahydro-2-oxo-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester with TFA, *p*-nitrophenyl

chloroformate, diisopropylethyl amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product. LC-MS: 471 (M^++1). 1H NMR ($CDCl_3$) δ 7.65 (d, 1H), 7.55 (s, 1H), 7.10 (d, 1H), 6.22 (d, 1H), 6.05 (s, 1H), 4.85 (s, 2H), 4.50 (m, 1H), 3.60 (m, 4H), 3.50 (m, 1H), 3.40 (m, 1H), 3.30 (m, 1H), 3.20 (m, 1H), 3.06 (m, 4H), 2.05 (m, 1H), 1.70-1.90 (m, 3H), 1.40 (m, 2H), 0.90 (m, 1H), 0.45 (m, 2H), 0.20 (m, 2H).

Example 199 :

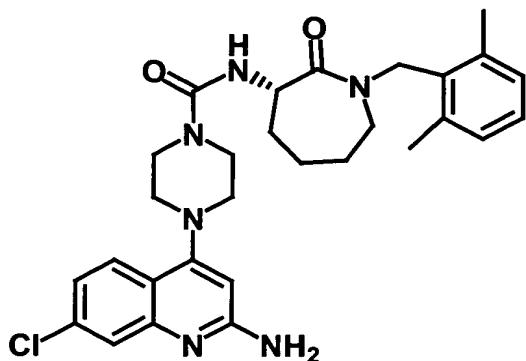
Preparation of 4-(2-Amino-7-chloro-4-quinoliny)-N-[(3S)-hexahydro-2-oxo-1-(phenylmethyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide



As described for example 78, (3*R*)-3-aminohexahydro-1-benzyl-2*H*-azepin-2-one obtained by de-protection of [(3*S*)-hexahydro-2-oxo-1-(phenylmethyl)-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester with TFA, *p*-nitrophenyl chloroformate, diisopropylethyl amine and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product. LC-MS 506 (M^++1). 1H NMR ($CDCl_3$) δ 7.70 (d, 1H), 7.58 (s, 1H), 7.30 (m, 5H), 7.15 (d, 1H), 6.35 (d, 1H), 6.12 (s, 1H), 4.90 (s, 2H), 4.64 (m, 1H), 4.50 (m, 2H), 3.65 (m, 4H), 3.50 (m, 1H), 3.22 (m, 1H), 3.10 (m, 4H), 2.10 (m, 1H), 1.70-2.00 (m, 3H), 1.50 (m, 1H), 1.20 (m, 1H).

Example 200 :

Preparation of 4-(2-Amino-7-chloro-4-quinoliny)-N-[(3*S*)-1-[(2,6-dimethylphenyl)methyl]hexahydro-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide



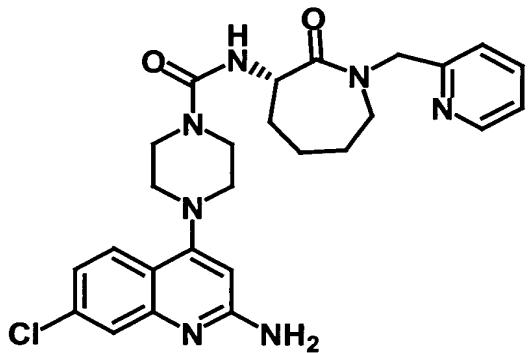
As described for example 78, (3*R*)-3-amino-1-[(2,6-dimethylphenyl)methyl]hexahydro-2*H*-azepin-2-one obtained by de-protection of

[(3*S*)-1-[(2,6-dimethylphenyl)methyl]hexahydro-2-oxo-1*H*-azepin-3-yl]-carbamic acid,

5 1,1-dimethylethyl ester with TFA, *p*-nitrophenyl chloroformate, diisopropylethyl amine and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product. LC-MS 535 (M^++1).

Example 201:

10 Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-*N*-(3*S*)-hexahydro-2-oxo-1-(2-pyridinylmethyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide

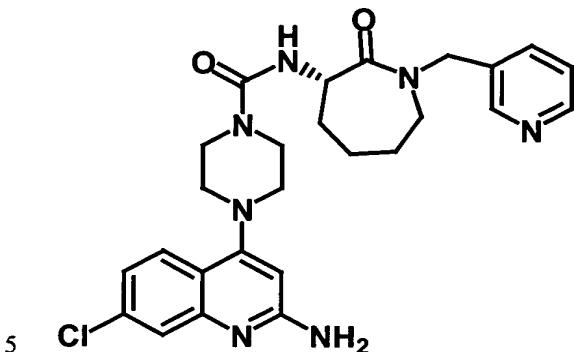


As described for example 78, (3*R*)-3-aminohexahydro-1-(2-pyridinylmethyl)-2*H*-azepin-2-one obtained by de-protection of [(3*S*)-hexahydro-2-oxo-1-(2-

15 pyridinylmethyl)-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester with TFA, *p*-nitrophenyl chloroformate, diisopropylethyl amine and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product. LC-MS 508 (M^++1). 1H NMR ($CDCl_3$) δ 8.50 (m, 1H), 7.60 (m, 3H), 7.15 (m, 3H), 6.20 (m, 1H), 6.05 (s, 1H), 4.80 (s, 2H), 4.70 (m, 2H), 4.60 (m, 1H), 3.60 (m, 4H), 3.50 (m, 1H), 3.40 (m, 1H), 3.05 (m, 4H),
20 2.10 (m, 1H), 1.60-1.90 (m, 3H), 1.50 (m, 1H), 1.30 (m, 1H).

Example 202 :

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[*(3S*)-hexahydro-2-oxo-1-(3-pyridinylmethyl)-1*H*-azepin-3-yl]-1-piperazin carboxamide

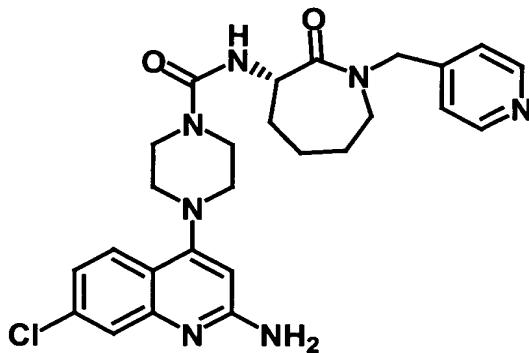


As described for example 78, (*3R*)-3-aminohexahydro-1-(3-pyridinylmethyl)-2*H*-azepin-2-one obtained by de-protection of [*(3S*)-hexahydro-2-oxo-1-(3-pyridinylmethyl)-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester with TFA, *p*-nitrophenyl chloroformate, diisopropylethyl amine and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product. LC-MS 508 (M^++1). 1H NMR ($CDCl_3$) δ 8.50 (m, 2H), 7.65 (d, 1H), 7.52 (m, 2H), 7.20 (m, 1H), 7.10 (d, 1H), 6.18 (d, 1H), 6.10 (s, 1H), 4.80 (s, 2H), 4.65 (m, 2H), 4.60 (m, 1H), 3.60 (m, 4H), 3.45 (m, 1H), 3.20 (m, 1H), 3.06 (m, 4H), 2.07 (m, 1H), 1.65-1.90 (m, 3H), 1.47 (m, 1H), 1.20 (m, 1H).

10

Example 203 :

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[*(3S*)-hexahydro-2-oxo-1-(4-pyridinylmethyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide



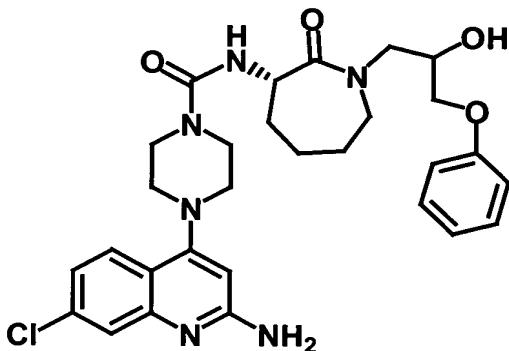
As described for example 78, (*3R*)-3-aminohexahydro-1-(4-pyridinylmethyl)-2*H*-azepin-2-one obtained by de-protection of [*(3S*)-hexahydro-2-oxo-1-(4-pyridinylmethyl)-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester with TFA, *p*-

20

nitrophenyl chloroformate, diisopropylethyl amine and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product. LC-MS 508 (M^++1).

Example 204 :

5 Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-hexahydro-1-(2-hydroxy-3-phenoxypropyl)-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide



*Preparation of [(3S)-hexahydro-1-(2-hydroxy-3-phenoxypropyl)-2-oxo-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester:* To a solution of [(3*R*)-hexahydro-2-oxo-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester (500 mg) in THF at -78 °C was

10 added LiHMDS (4.4 mL, 1.0 M in THF). After addition, the reaction mixture was allowed to warm to rt. After 30 min at rt, the reaction mixture was re-cooled to -78 °C, then 2-(phenoxyethyl)-oxirane (0.6 mL) was introduced. The reaction was kept at rt for additional 1 h, and was quenched with NH₄Cl (sat., 20 mL). The reaction mixture
15 was extracted with EtOAc, washed with brine, and dried over Na₂SO₄. Concentration *in vacuo* afforded the crude product which was used directly for next step reaction without further purification.

In a similar manner, [(3*S*)-hexahydro-1-(2-hydroxypropyl)-2-oxo-1*H*-azepin-3-

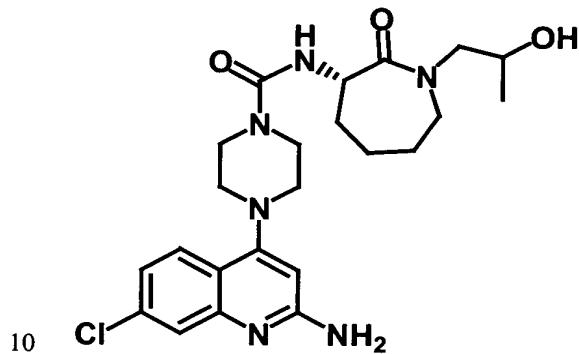
20 yl]- carbamic acid, 1,1-dimethylethyl ester was prepared.

*Preparation of 4-(2-amino-7-chloro-4-quinolinyl)-N-[(3*S*)-hexahydro-1-(2-hydroxy-3-phenoxypropyl)-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide:* As described for example 78, (3*R*)-3-aminohexahydro-1-(2-hydroxy-3-phenoxypropyl)-2*H*-azepin-2-one
25 obtained by de-protection of [(3*S*)-hexahydro-1-(2-hydroxy-3-phenoxypropyl)-2-oxo-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester with TFA, 4-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine

are reacted to afford the product as a light yellow solid. LC-MS: 566 (M^++1). 1H NMR (CDCl₃) δ 1.4-1.65 (m, 2H), 1.7-2.24 (m, 4H), 3.05-3.2 (m, 4H), 3.34-3.5 (m, 2H), 3.56-3.74 (m, 6H), 3.75-4.04 (m, 3H), 4.13-4.28 (m, 1H), 4.6-4.75 (m, 1H), 4.8-5.0 (s, 2H), 6.1-6.25 (m, 2H), 6.88-6.94 (m, 2H), 6.95-7.05 (m, 1H), 7.14-7.2 (m, 1H), 7.26-7.33 (m, 2H), 7.6-7.63 (d, 1H), 7.7-7.75 (d, 1H), 8.56-8.74 (m, 1H).

Example 205 :

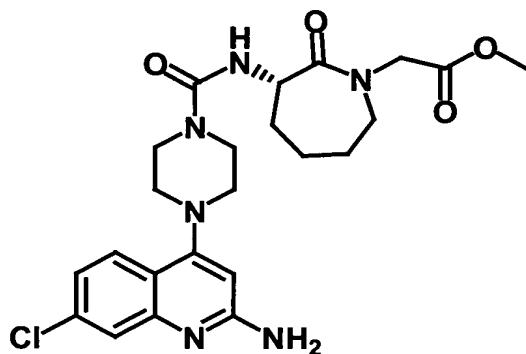
Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-hexahydro-1-(2-hydroxypropyl)-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide



As described for example 78, (3*R*)-3-aminohexahydro-1-(2-hydroxypropyl)-2*H*-azepin-2-one obtained by de-protection of [(3*S*)-hexahydro-1-(2-hydroxypropyl)-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester with TFA, 4-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a light yellow solid. LC-MS: 474 (M^++1). 1H NMR (CDCl₃) δ 0.75-1.0 (m, 1H), 1.18-1.3 (m, 3H), 1.3-1.6 (m, 2H), 1.7-2.4 (m, 4H), 3.05-3.2 (m, 4H), 3.2-3.36 (m, 3H), 3.6-3.8 (m, 4H), 3.9-4.1 (m, 1H), 4.45-4.7 (m, 1H), 4.96-5.1 (s, 2H), 6.1-6.3 (m, 3H), 7.1-7.2 (m, 1H), 7.6 (m, 1H), 7.7-7.8 (m, 1H).

Example 206 :

Preparation of (3*S*)-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-acetic acid, methyl ester



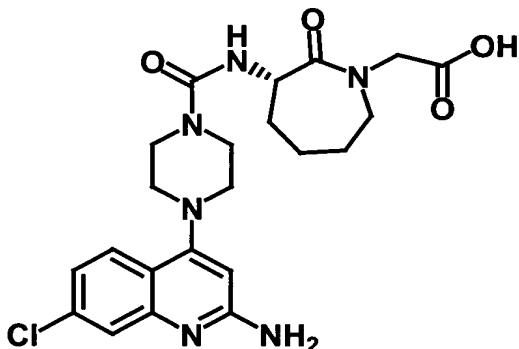
As described for example 78, (3S)-3-aminohexahydro-2-oxo-1H-azepine-1-acetic acid, methyl ester obtained by de-protection of (3S)-3-[(1,1-dimethylethoxy)carbonyl]amino]hexahydro-2-oxo-1H-azepine-1-aceticacid, methyl

5 ester with TFA, *p*-nitrophenyl chloroformate, diisopropylethyl amine and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product. LC-MS 489 ($M^{++}1$). 1H NMR ($CDCl_3$) δ 7.70 (d, 1H), 7.60 (s, 1H), 7.18 (d, 1H), 6.15 (m, 2H), 4.80 (s, 2H), 4.65 (m, 1H), 4.20 (s, 2H), 3.75 (m, 4H), 3.65 (m, 4H), 3.20 (m, 1H), 3.10 (m, 4 H), 2.10 (m, 1H), 1.80-2.05 (m, 3H), 1.60 (m, 2H).

10

Example 207 :

Preparation of (3S)-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1H-azepine-1-acetic acid



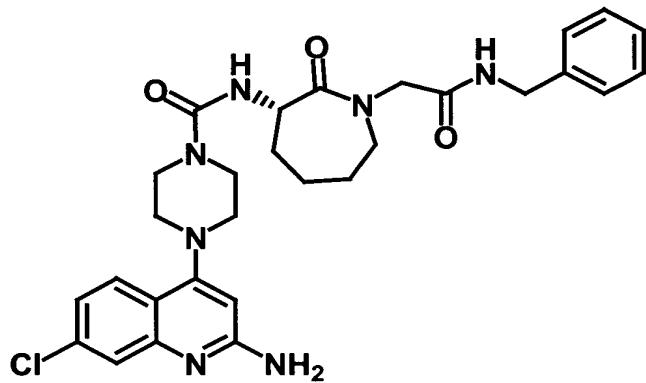
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To a stirred solution of (3S)-[[[4-(2-amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1H-azepine-1-acetic acid, methyl ester (300mg, 0.613 mmol) in THF (3.0 mL) was added 2N NaOH solution (3 mL) at rt. After 30 min, THF was evaporated *in vacuo*, and the resulting residue was acidified to pH 4-5. The reaction mixture was extracted with ethyl acetate, washed with brine, and dried over Na_2SO_4 . The product (15 mg) was obtained after concentration *in vacuo*.

LC-MS: 475 (M^++1). 1H NMR (CD_3OD) δ 7.82 (d, 1H), 7.50 (s, 1H), 7.38 (d, 1H), 6.25 (s, 1H), 4.58 (d, 1H), 4.10 (m, 2H), 3.80 (m, 1H), 3.65 (m, 4H), 3.30 (m, 5H), 1.60-1.96 (m, 6H).

5 **Example 208 :**

Preparation of (3S)-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-N-(phenylmethyl)-1*H*-azepine-1-acetamide



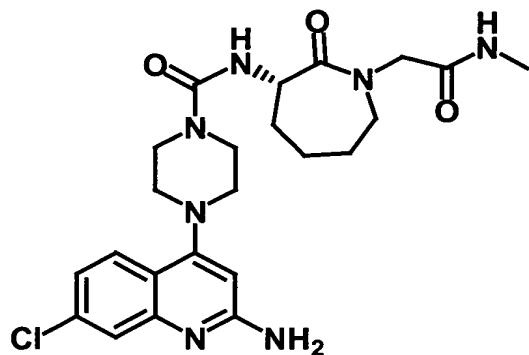
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The mixtures of (3S)-[[[4-(2-amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-acetic acid (50.0mg, 0.1 mmol), benzyl amine (0.2 mL), HATU (50 mg), and diisopropylethyl amine (0.3 mL) in DMF (5 mL) was stirred at rt for 2h. The reaction mixture was diluted with ethyl acetate (100 mL), washed with $NaHCO_3$ (sat.) and water, dried over Na_2SO_4 .
Concentration *in vacuo* followed by flash chromatography afforded the title product.
LC-MS: 564 (M^++1). 1H NMR (CD_3OD) δ 7.82 (d, 1H), 7.45 (s, 1H), 7.25 (m, 4H), 7.20 (m, 2H), 6.30 (s, 1H), 4.65(d, 1H), 4.40 (s, 2H), 4.15 (m, 2H), 3.75 (m, 1H), 3.65(m, 4H), 3.30 (m, 1H), 3.10 (m, 4H), 1.96 (m, 2H), 1.80 (m, 2H), 1.60 (m, 2H).

20

Example 209 :

Preparation of (3S)-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-N-methyl-2-oxo-1*H*-azepine-1-acetamide

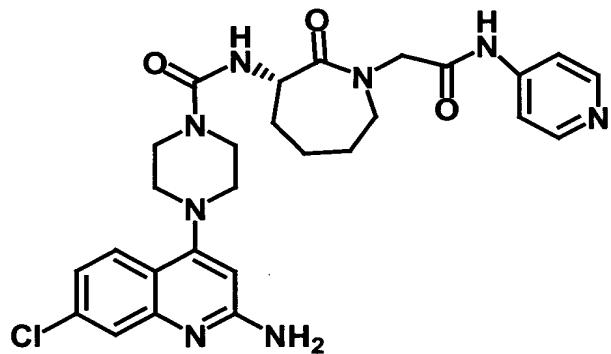


As described as example 208, (3S)-[[[4-(2-amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-acetic acid, methyl amine, HATU, and diisopropylethylamine are reacted to afford the product. LC-MS 488

5 (M⁺+1). ¹H NMR (CD₃OD) δ 7.86 (d, 1H), 7.50 (s, 1H), 7.30 (d, 1H), 6.30 (s, 1H), 4.65 (m, 1H), 4.00 (m, 2H), 3.70(m, 5H), 3.30 (m, 5H), 2.75 (s, 3H), 1.98 (m, 2H), 1.80 (m, 2H), 1.60 (m, 2H).

Example 210 :

10 Preparation of (3S)-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-N-(4-pyridinyl)-1*H*-azepine-1-acetamide

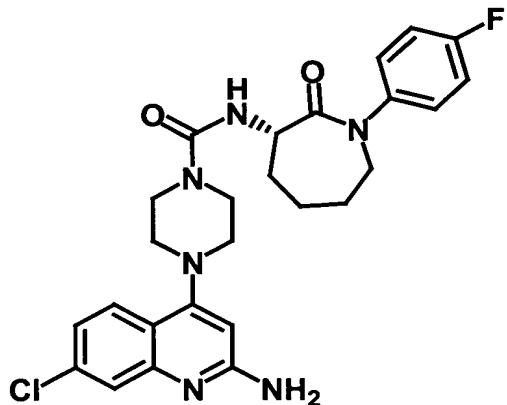


15 As described as example 208, (3S)-[[[4-(2-amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-acetic acid, 4-aminopyridine, HATU, and diisopropylethylamine are reacted to afford the product. LC-MS: 551 (M⁺+1). ¹H NMR (CD₃OD) δ 1.60-1.80 (m, 4H), 1.90 (m, 2H), 3.10 (m, 4H), 3.30 (m, 1H), 3.60 (m, 4H), 4.20 (m, 2H), 4.60 (d, 1H), 6.24 (s, 1H), 7.08 (d, 1H), 7.39 (s, 1H), 7.55 (m, 2H), 7.75 (d, 2H), 8.30 (m, 2H).

20

Example 211 :

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3R)-1-(4-fluorophenyl)hexahydro-2-oxo-1H-azepin-3-yl]-1-piperazinecarboxamide

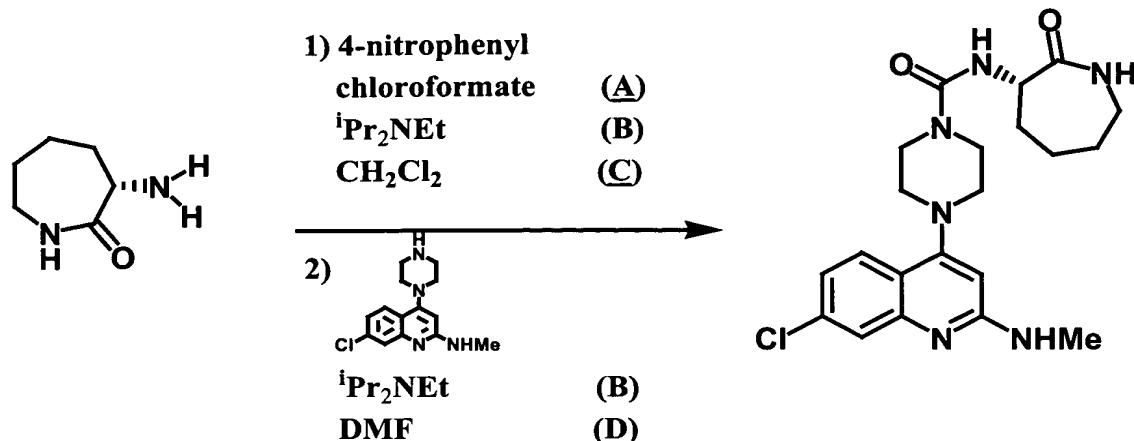


Preparation of [(3S)-1-(4-fluorophenyl)hexahydro-2-oxo-1H-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester (According to *Tetrahedron Lett.*, **1998**, 39, 2933-2936):
 5 The mixture of [(3R)-hexahydro-2-oxo-1H-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester 400 mg), 4-fluorophenylboronic acid (200 mg), Et₃N (0.3 mL), and Cul (150 mg) in CH₂Cl₂ (5 mL) are mixture at rt for 4 days. The reaction mixture was diluted with brine, extracted with EtOAc, and dried. Concentration followed by flash
 10 chromatograph afforded the product (180 mg).

Preparation of 4-(2-amino-7-chloro-4-quinolinyl)-N-[(3R)-1-(4-fluorophenyl)hexahydro-2-oxo-1H-azepin-3-yl]-1-piperazinecarboxamide: As described for example 78, (3S)-3-amino-1-(4-fluorophenyl)hexahydro-2H-azepin-2-one obtained by the de-protection of
 15 [(3S)-1-(4-fluorophenyl)hexahydro-2-oxo-1H-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester with TFA, 4-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a light yellow solid. LC-MS: 511 (M⁺+1). ¹H NMR (CDCl₃) δ 1.62 (m, 2H), 1.83-2.31 (m, 4H), 3.14 (m, 4H), 3.62 (m 4H), 3.95 (dd, 1H), 4.81 (m, 1H), 6.17 (s, 1H), 6.22 (d, 1H), 7.06
 20 (m, 2H), 7.18 (m, 2H), 7.6 (s, 1H), 7.65 (d, 1H).

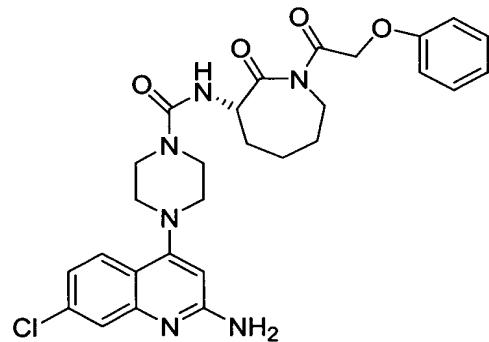
Example 212 :

Preparation of 4-[7-Chloro-2-(methylamino)-4-quinolinyl]-N-[(3S)-hexahydro-2-oxo-1H-azepin-3-yl]-1-piperazinecarboxamide



As described for example 78, (3*S*)-3-aminohexahydro-2*H*-azepin-2-one, 4-nitrophenyl
 5 chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine
 are reacted to afford the product as a light yellow solid. LC-MS: 431(M⁺+1). ¹H NMR
 (CDCl₃): δ 1.34~1.60 (m, 2H), 1.80~1.92 (m, 2H), 1.98~2.08 (m, 1H), 2.10~2.18 (m,
 1H), 3.06 (d, 3H), 3.12 (br.s, 4H), 3.20~3.38 (m, 2H), 3.67 (br.s, 4H), 4.52 (br. dd, 1H),
 5.98~6.06 (m, 2H), 6.12 (d, 1H), 7.12 (dd, 1H), 7.67 (d, 1H), 7.70 (d, 1H).

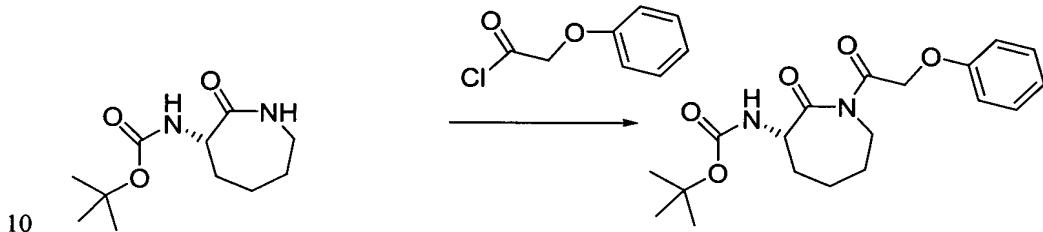
10

Example 213 :**Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3*S*)-hexahydro-2-oxo-1-(2-phenoxyacetyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide**

15 Preparation of [(3*S*)-hexahydro-2-oxo-1-(phenoxyacetyl)-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester (A. D. Borthwick, et al. *J. Med. Chem.*, **2000**, *43*, 4452-4464): To a stirred solution of [(3*S*)-hexahydro-2-oxo-1*H*-azepin-3-yl]-carbamic acid-1,1-dimethylethyl ester (500 mg, 2.2 mmol) in THF (10 mL) at -78 °C was added LiHDS (3 mL, 3 mmol, 1.0 M in THF). After addition, dry-ice bath was removed, and

the reaction was stirred at room temperature for about 30 min. The reaction was re-cooled to -78 °C, and phenoxyacetyl chloride (0.91 mL, 6.58 mmol) was added. The reaction was allowed to warm to rt over 2 h, and was quenched with NH₄Cl (sat., 10 mL). The reaction mixture was extracted with EtOAc (3x30 mL), washed with brine

5 (15 mL), and dried over Na₂SO₄. Concentration *in vacuo* followed by flash chromatography (hexane: EtOAc/ 5:1) afforded the title product. ¹H NMR (CDCl₃) δ 1.38 (m, 1H), 1.42 (s, 9H), 1.63 (m, 1H), 1.77-1.98 (m, 3H), 2.18 (m, 1H), 3.21 (m, 1H), 4.67 (m, 1H), 4.83 (m, 1H), 5.2 (m, 1H), 5.61 (m, 1H), 6.9 (m, 3H), 7.28 (m, 2H).



The following intermediates were prepared in a similar manner:

[(3S)-1-acetylhexahydro-2-oxo-1H-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester

[(3S)-1-(cyclopropylcarbonyl)hexahydro-2-oxo-1H-azepin-3-yl]-carbamic acid,

15 1,1-dimethylethyl ester

[(3S)-1-benzoylhexahydro-2-oxo-1H-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester

[(3S)-1-(ethylsulfonyl)hexahydro-2-oxo-1H-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester

20 [(3S)-hexahydro-2-oxo-1-(phenylsulfonyl)-1H-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester

[(3S)-1-(cyclohexylcarbonyl)hexahydro-2-oxo-1H-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester

25 [(3S)-hexahydro-2-oxo-1-[[[phenylsulfonyl]amino]carbonyl]-1H-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester

[(3S)-hexahydro-2-oxo-1-(phenylacetyl)-1H-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester

[(3R)-1-formylhexahydro-2-oxo-1H-azepin-3-yl]-carbamic acid-1,1-dimethylethyl ester

[(3R)-1-(2,2-dimethyl-1-oxopropyl)hexahydro-2-oxo-1H-azepin-3-yl]- carbamic acid-1,1-dimethylethyl ester

[(3R)-hexahydro-1-(methylsulfonyl)-2-oxo-1H-azepin-3-yl]-carbamic acid-1,1-dimethylethyl ester

5 3R)-3-[[[(1,1-dimethylethoxy)carbonyl]amino]hexahydro-2-oxo-1H-azepine-1-carboxylic acid, methyl ester

(3R)-3-[[[(1,1-dimethylethoxy)carbonyl]amino]hexahydro-2-oxo-1H-azepine-1-carboxylic acid-ethyl ester

(3R)-3-[[[(1,1-dimethylethoxy)carbonyl]amino]hexahydro-2-oxo-1H-azepine-1-carboxylic acid, 2,2,2-trifluoroethyl ester

10 (3S)-3-[[[(1,1-dimethylethoxy)carbonyl]amino]hexahydro-1H-azepine-1-carboxylic acid 2-oxo-, 2-propenyl ester

(3R)-3-[[[(1,1-dimethylethoxy)carbonyl]amino]hexahydro-2-oxo-1H-azepine-1-carboxylic acid, phenyl ester

15 (3R)-3-[[[(1,1-dimethylethoxy)carbonyl]amino]hexahydro-2-oxo-1H-azepine-1-carboxylic acid-4-fluorophenyl ester

(3S)-3-[[[(1,1-dimethylethoxy)carbonyl]amino]hexahydro-2-oxo-1H-azepine-1-carboxylic acid, phenylmethyl ester

[(3S)-1-cyanohexahydro-2-oxo-1H-azepin-3-yl]-carbamic-acid-1,1-dimethylethyl ester:

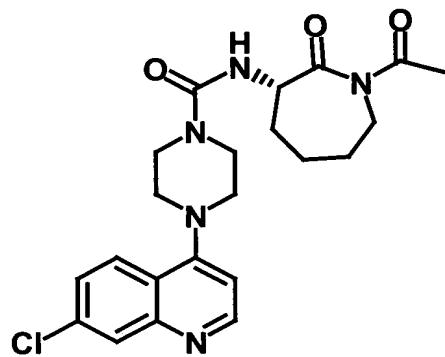
Preparation of 4-(2-amino-7-chloro-4-quinolinyl)-N-[(3S)-hexahydro-2-oxo-1-(2-phenoxyacetyl)-1H-azepin-3-yl]-1-piperazinecarboxamide: To a mixture of 3-aminohexahydro-1-(phenoxyacetyl)-2H-azepin-2-one (~0.43mmol) [obtained by the de-protection of de-protection of [(3S)-hexahydro-2-oxo-1-(phenoxyacetyl)-1H-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester with TFA] in CH₂Cl₂ (2mL) and NaHCO₃ (sat., 2 mL) was added triphosgene (42mg, 0.14mmol) in at 0 °C. After 15 min, the mixture was diluted with water and extracted with minimum CH₂Cl₂, then dried over Na₂SO₄, which was used directly in the next step. To this solution was added a hot DMSO (3 mL) solution of 7-chloro-4-(1-piperazinyl)-2-quinolinamine followed by iPr₂NEt (0.1mL, 0.57mmol,). After 1.5 h at rt, the mixture was poured into ice-water. The reaction mixture was extracted with EtOAc, washed with brine, and dried over Na₂SO₄. Concentration in vacuo followed by preparative TLC (CH₂Cl₂-MeOH-

$\text{Et}_3\text{N}, 9:1:1\%$) afforded the title product as a light yellow solid. LC-MS: 551 (M^++1). ^1H NMR (CDCl_3): δ 1.32~2.24 (m, 6H), 3.14~3.30 (m, 5H), 3.70 (m, 4H), 4.76~4.86 (m, 3H), 4.94 (m, 1H), 5.16 (s, 2H), 5.78 (br.d, 1H), 6.16 (s, 1H), 6.92 (m, 2H), 6.99 (m, 1H), 7.18 (dd, 1H), 7.26~7.32 (m, 3H), 7.62 (d, 1H), 7.73 (d, 1H).

5

Example 214 :

Preparation of 4-(7-Chloro-4-quinolinyl)-*N*-[(3*S*)-1-acetylhexahydro-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide:

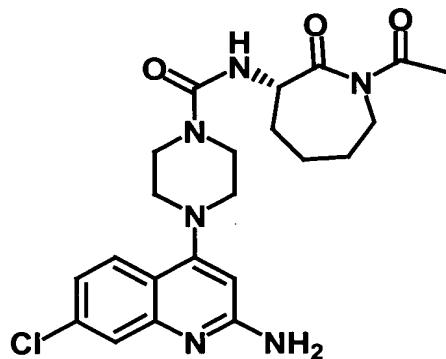


10 As described for example 78, (3*R*)-1-acetyl-3-amino hexahydro-2*H*-azepin-2-one obtained by de-protection of [(3*S*)-1-acetylhexahydro-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester with TFA, diisopropyl(ethyl)amine, diisopropyl(ethyl)amine, and 7-chloro-4-(piperazin-1-yl)quinoline are reacted to give the product. LC-MS: 444 (M^++1). ^1H NMR (CDCl_3) δ 8.68 (d, 1H), 8.00 (s, 1H), 7.90 (d, 1H), 7.40 (d, 1H), 6.80 (d, 1H), 5.80 (d, 1H), 4.83 (m, 1H), 4.70 (m, 1H), 3.65 (m, 4H), 3.18 (m, 5H), 2.50 (s, 3H), 2.10 (m, 1H), 1.90 (m, 3H), 1.50 (m, 1H), 1.30 (m, 1H).

15

Example 215 :

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)- N-[(3S)-1-acetylhexahydro-2-oxo-1H-azepin-3-yl]-1-piperazinecarboxamide



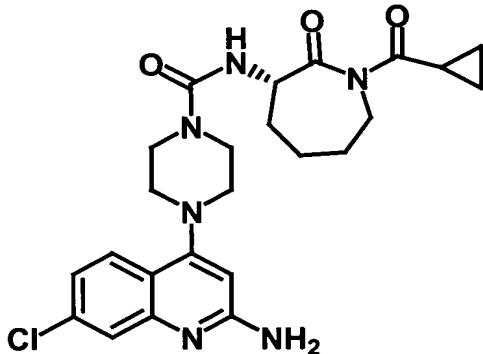
5

As described for example 78, (3R)-1-acetyl-3-aminohexahydro-2H-azepin-2-one obtained by de-protection of [(3S)-1-acetylhexahydro-2-oxo-1H-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester with TFA, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to give the product. LC-MS: 459 (M^++1). 1H

10 $NMR (CDCl_3) \delta 7.65 (d, 1H), 7.55 (s, 1H), 7.10 (d, 1H), 6.10 (s, 1H), 5.80 (m, 1H), 5.15 (s, 2H), 4.85 (m, 1H), 4.70 (m, 1H), 3.60 (m, 4H), 3.10 (m, 5H), 2.45 (s, 3H), 2.10 (m, 1H), 1.85 (m, 3H), 1.50 (m, 1H), 1.30 (m, 1H).$

Example 216 :

15 **Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-1-(cyclopropylcarbonyl)hexahydro-2-oxo-1H-azepin-3-yl]-1-piperazinecarboxamide:**

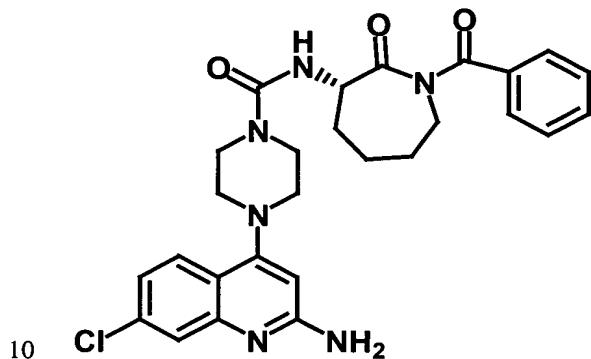


As described for example 78, (3R)-3-amino-1-(cyclopropylcarbonyl)hexahydro-2H-azepin-2-one obtained by de-protection of [(3S)-1-(cyclopropylcarbonyl)hexahydro-2-oxo-1H-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester with TFA,

diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to give the product. LC-MS: 485 (M^++1). ^1H NMR (CDCl_3) δ 7.65 (d, 1H), 7.58 (s, 1H), 7.12 (d, 1H), 6.06 (s, 1H), 5.86 (d, 1H), 4.95 (s, 2H), 4.80 (m, 1H), 4.58 (m, 1H), 3.60 (m, 4H), 3.10 (m, 5H), 2.70 (m, 1H), 2.10 (m, 1H), 1.70-1.90 (m, 3H), 1.60 (m, 1H),
5 1.30 (m, 1H), 1.10 (m, 2H), 0.90 (m, 2H).

Example 217 :

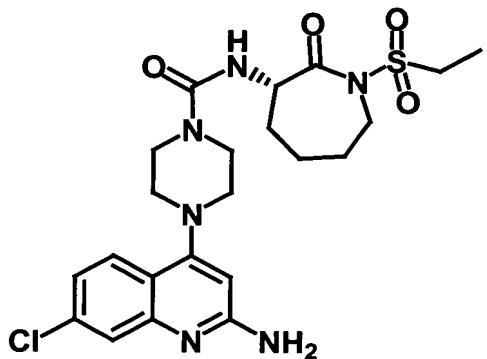
Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-1-benzoylhexahydro-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide



As described for 78, (3*R*)-3-amino-1-benzoylhexahydro-2*H*-azepin-2-one obtained from de-protection of [(3*S*)-1-benzoylhexahydro-2-oxo-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester with TFA, 4-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product. LC-MS: 521 (M^++1). ^1H NMR (CDCl_3) δ 7.75 (d, 1H), 7.60 (m, 3H), 7.50 (m, 1H), 7.43 (m, 2H), 7.20 (d, 1H), 6.10 (s, 1H), 5.75 (d, 1H), 4.85 (m, 1H), 4.75 (s, 2H), 4.55 (m, 1H), 3.60 (m, 4H), 3.50 (m, 1H), 3.10 (m, 4H), 2.25 (m, 1H), 2.10 (m, 3H), 1.70 (m, 2H).
10
15

Example 218 :

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3*S*)-1-(ethylsulfonyl)hexahydro-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide

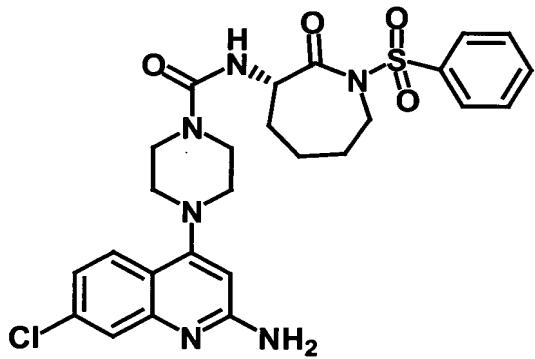


As described for 78, (3*R*)-3-amino-1-(ethylsulfonyl)hexahydro-2*H*-azepin-2-one obtained from the de-protection of [(3*S*)-1-(ethylsulfonyl)hexahydro-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester with TFA, 4-nitrophenyl chloroformate,

5 diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product. LC-MS: 509 (M^++1). ^1H NMR (CDCl_3) δ 7.75 (d, 1H), 7.60 (s, 1H), 7.20 (d, 1H), 6.15 (s, 1H), 5.80 (d, 1H), 4.80 (m, 3H), 4.40 (m, 1H), 3.65 (m, 4H), 3.45-3.60 (m, 2H), 3.16 (m, 4H), 2.18 (m, 1H), 2.00 (m, 2H), 1.70 (m, 4H), 1.35 (t, 3H).

10 **Example 219 :**

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3*S*)-hexahydro-2-oxo-1-(phenylsulfonyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide

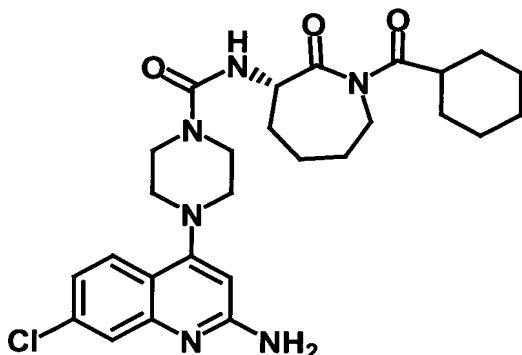


As described for 78, (3*R*)-3-aminohexahydro-1-(phenylsulfonyl)-2*H*-azepin-2-one

15 obtained from de-protection of [(3*S*)-hexahydro-2-oxo-1-(phenylsulfonyl)-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester with TFA, 4-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product. LC-MS: 557 (M^++1).

20 **Example 220 :**

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-1-(cyclohexylcarbonyl)hexahydro-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide

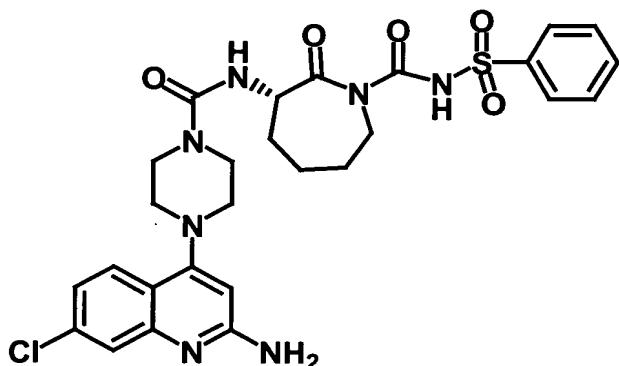


5 As described for 78, (3*R*)-3-amino-1-(cyclohexylcarbonyl)hexahydro-2*H*-azepin-2-one obtained from de-protection of [(3*S*)-1-(cyclohexylcarbonyl)hexahydro-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester with TFA, 4-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product. LC-MS: 527 (M^++1). 1H NMR ($CDCl_3$) δ 7.75 (d, 1H), 7.60 (s, 1H), 7.18 (d, 1H), 6.15 (s, 1H), 5.98 (d, 1H), 4.90 (m, 3H), 4.65 (m, 1H), 3.70 (m, 4H), 3.75 (m, 1H), 3.12 (m, 3H), 2.20 (m, 1H), 1.60-1.95 (m, 10H), 1.20-1.40 (m, 6H).

10

15 **Example 221 :**

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3*S*)-hexahydro-2-oxo-1-[[*(phenylsulfonyl)amino*]carbonyl]-1*H*-azepin-3-yl]-1-piperazinecarboxamide



As described for 78, *N*-[(3*R*)-3-aminohexahydro-2-oxo-1*H*-azepin-1-yl]carbonyl]-benzenesulfonamide obtained from de-protection of [(3*S*)-hexahydro-2-oxo-1-

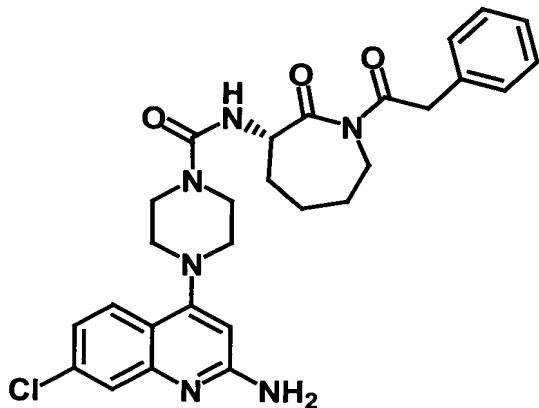
20

[[(phenylsulfonyl)amino]carbonyl]-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester with TFA, 4-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product. LC-MS: 600 (M^++1).

¹H NMR ($CDCl_3$) δ 8.10 (m, 2H), 7.80 (m, 2H), 7.50 (m, 4H), 7.40 (m, 1H), 6.15 (s, 1H),
 5 5.25 (m, 1H), 4.60 (m, 1H), 3.60-3.80 (m, 5H), 3.30 (m, 5H), 2.10 (m, 1H), 2.00 (m, 1H), 1.80 (m, 2H), 1.60 (m, 2H).

Example 222 :

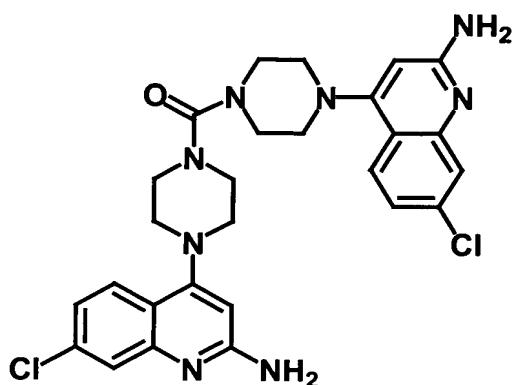
10 **Preparation of 4-(2-Amino-7-chloro-4-quinoliny)-N-[*(3S*)-hexahydro-2-oxo-1-(2-phenylacetyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide**



As described for example 213, [(3*S*)-hexahydro-2-oxo-1-(phenylacetyl)-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester, TFA, triphosgene, NaHCO₃ (sat.), and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product
 15 as a light yellow solid. LC-MS: 535 (M^++1). ¹H NMR ($CDCl_3$, 400MHz): δ 1.20~1.50 (m, 2H), 1.80~2.20 (m, 4H), 3.10~3.20 (m, 5H), 3.60~3.70 (m, 4H), 4.28 (AB, 2H), 4.75 (dd, 1H), 4.80~4.95 (m, 3H), 5.85 (d, 1H), 6.16 (s, 1H), 7.18 (dd, 1H), 7.22~7.36 (m, 5H), 7.62 (d, 1H), 7.72 (d, 1H).

20

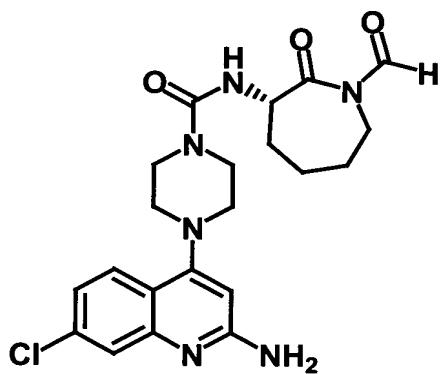
4-[4-[[4-(2-Amino-7-chloro-4-quinoliny)-1-piperazinyl]carbonyl]-1-piperazinyl]-7-chloro-2-quinolinamine



5 The title compound was obtained as a product in example 222. LC-MS: 551(M⁺+1). ¹H NMR (DMSO-d₆-TFA): δ 2.95 (br.s, 8H), 3.13(br.s, 8H), 5.98 (s, 2H), 7.06 (dd, 2H), 7.29 (d, 2H), 7.54 (d, 2H), 8.92 (br.s, 4H).

10 **Example 223 :**

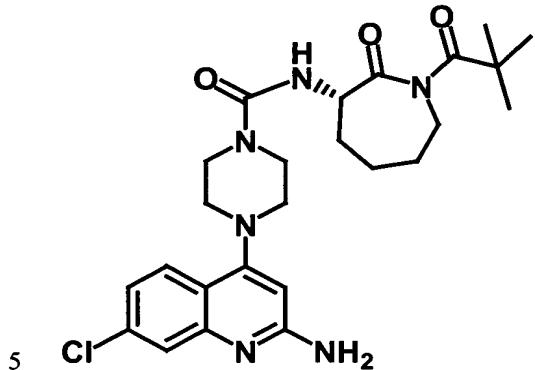
Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-1-formylhexahydro-2-oxo-1H-azepin-3-yl]-1-piperazinecarboxamide



As described for example 213, [(3R)-1-formylhexahydro-2-oxo-1H-azepin-3-yl]-15 carbamic acid-1,1-dimethylethyl ester, TFA, triphosgene, NaHCO₃ (sat.), and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a light yellow solid. LC-MS: 445 (M⁺+1). ¹H NMR (CDCl₃): δ 1.30 (m, 1H), 1.65 (m, 1H), 1.80~2.30 (m, 4H), 3.09 (dd, 1H), 3.14 (m, 4H), 3.68 (m, 4H), 4.62 (br.dd, 1H), 4.82 (ddd, 1H), 4.90 (br.s, 2H), 5.84 (d, 1H), 6.16 (s, 1H), 7.18 (dd, 1H), 7.62 (d, 1H), 7.72 (d, 1H), 20 9.47 (s, 1H).

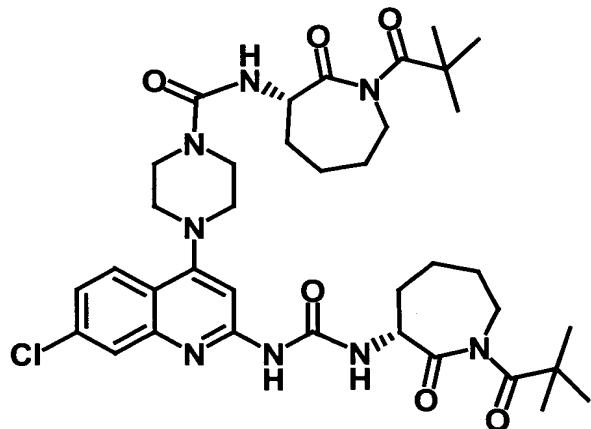
Example 224 :

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-1-(2,2-dimethyl-1-oxopropyl)hexahydro-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide



As described for example 213, [(3*R*)-1-(2,2-dimethyl-1-oxopropyl)hexahydro-2-oxo-1*H*-azepin-3-yl]- carbamic acid-1,1-dimethylethyl ester, TFA, triphosgene, NaHCO₃ (sat.), and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a light yellow solid. LC-MS: 501(M⁺+1). ¹H NMR (CDCl₃) δ 1.30 (s, 9H), 1.50~1.70 (m, 1H), 1.80~2.08 (m, 3H), 2.10~2.20 (m, 1H), 3.05~3.20 (m, 4H), 3.45 (dd, 1H), 3.60~3.74 (m, 4H), 3.80 (br.dd, 1H), 4.66 (m, 1H), 4.94 (br.s, 2H), 5.96 (d, 1H), 6.15 (s, 1H), 7.16 (dd, 1H), 7.60 (d, 1H), 7.71 (d, 1H).

15 **4-[7-Chloro-2-[[[[3*R*)-1-(2,2-dimethyl-1-oxopropyl)hexahydro-2-oxo-1*H*-azepin-3-yl]amino]carbonyl]amino]-4-quinolinyl]-N-[(3*S*)-1-(2,2-dimethyl-1-oxopropyl)hexahydro-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide**

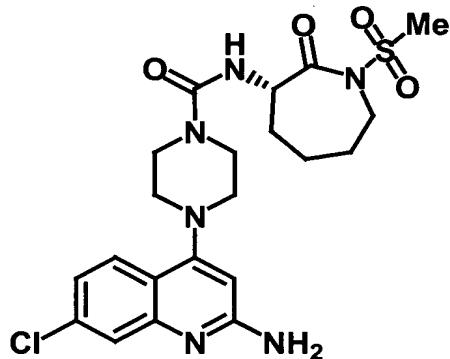


The title compound was obtained as a product in example 224. LC-MS: 739 (M^++1). 1H NMR (DMSO-d₆-TFA) δ 1.10~2.35 (m, 30H), 3.10~3.90 (m, 12H), 4.65 (m, 1H), 4.70 (m, 1H), 5.95 (d, 1H), 6.30 (s, 1H), 7.30 (dd, 1H), 7.60 (br.s, 1H), 7.80 (d, 1H), 7.95 (s, 1H).

5

Example 225 :

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-hexahydro-1-(methylsulfonyl)-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide



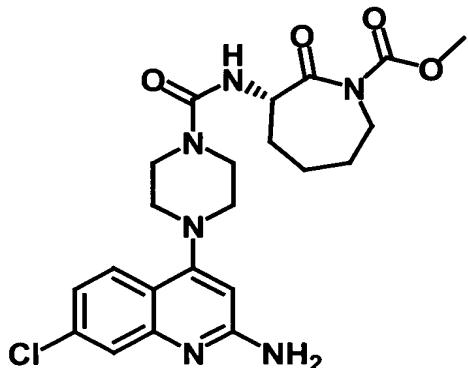
10

As described for example 213, [(3*R*)-hexahydro-1-(methylsulfonyl)-2-oxo-1*H*-azepin-3-yl]-carbamic acid-1,1-dimethylethyl ester, TFA, triphosgene, NaHCO₃ (sat.), and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a light yellow solid. LC-MS: 495 (M^++1). 1H NMR (CDCl₃) δ 1.52 (m, 1H), 1.70 (m, 1H), 1.86 (m, 1H), 1.95~2.05 (m, 2H), 2.15 (m, 1H), 3.09 (m, 4H), 3.45 (s, 3H), 3.48 (m, 1H), 3.65 (m, 4H), 4.45 (m, 1H), 4.81 (m, 1H), 5.17 (br.s, 2H), 5.95 (d, 1H), 6.15 (s, 1H), 7.15 (dd, 1H), 7.60 (d, 1H), 7.69 (d, 1H).

15

Example 226 :

20 **Preparation of (3*S*)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-carboxylic acid, methyl ester**

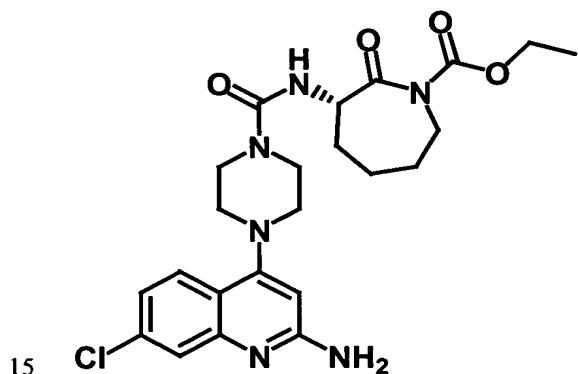


As described for example 213, (3*R*)-3-[(1,1-dimethylethoxy)carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-carboxylic acid-methyl ester, TFA, triphosgene, NaHCO₃ (sat.), and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the title product as a light yellow solid. LC-MS: 475 (M⁺+1). ¹H NMR (CDCl₃): δ 1.45~2.05 (m, 5H), 2.15 (m, 1H), 3.12 (m, 4H), 3.42 (dd, 1H), 3.66 (m, 4H), 3.90 (s, 3H), 4.45 (m, 1H), 4.75~4.85 (m, 3H), 5.95 (d, 1H), 6.15 (s, 1H), 7.19 (dd, 1H), 7.62 (d, 1H), 7.74 (d, 1H).

10

Example 227 :

Preparation of (3*S*)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-carboxylic acid-ethyl ester



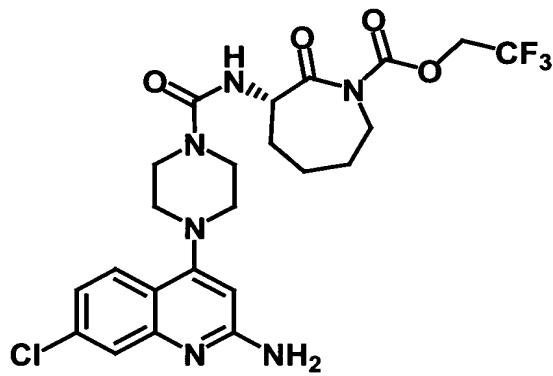
As described for example 213, (3*R*)-3-[(1,1-dimethylethoxy)carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-carboxylic acid, ethyl ester, TFA, triphosgene, NaHCO₃ (sat.), and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the title product as a light yellow solid. LC-MS:

489 (M^++1). 1H NMR ($CDCl_3$): δ 1.37 (t, 3H), 1.44~2.02 (m, 5H), 2.12~2.20 (m, 1H), 3.13 (m, 4H), 3.41 (dd, 1H), 3.67 (m, 4H), 4.35 (m, 2H), 4.42 (m, 1H), 4.74~4.86 (m, 3H), 6.00 (d, 1H), 6.16 (s, 1H), 7.18 (dd, 1H), 7.62 (d, 1H), 7.73 (d, 1H).

5 Example 228 :

Preparation of (3*S*)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-carboxylic acid,2,2,2-trifluoroethyl ester

10

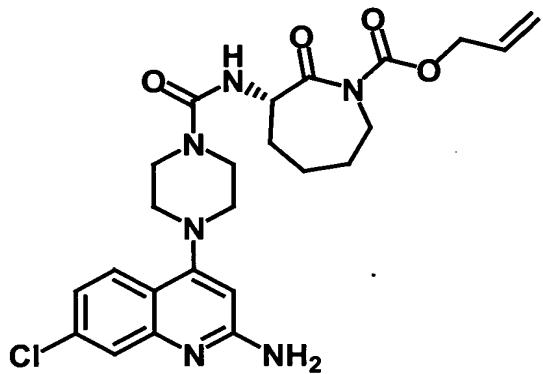


As described for example 213, (3*R*)-3-[(1,1-

15 dimethylethoxy)carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-carboxylic acid, 2,2,2-trifluoroethyl ester, TFA, triphosgene, $NaHCO_3$ (sat.), and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a white solid. LC-MS: 543 (M^++1).
 1H NMR (DMSO+TFA): δ 1.32~1.45 (m, 1H), 1.56~1.90 (m, 5H), 3.26 (m, 4H), 3.54~3.66 (m, 5H), 4.15 (m, 1H), 4.68 (br.d, 1H), 4.83 (m, 2H), 6.32 (s, 1H), 7.42 (dd, 1H), 7.60 (d, 1H), 7.90 (d, 1H).

20 Example 229 :

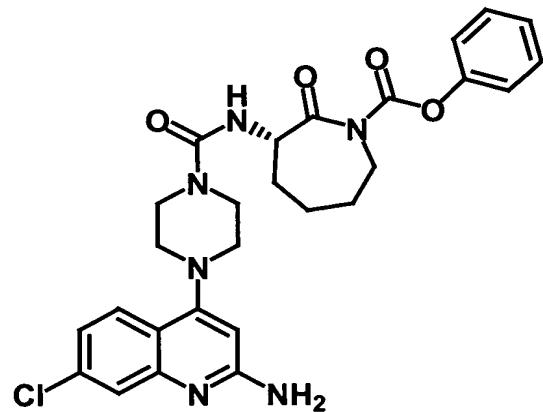
Preparation of (3*S*)-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-carboxylic acid, 2-propenyl ester



As described for example 213, (3*S*)-3-[(1,1-dimethylethoxy)carbonyl]amino]hexahydro-1*H*-azepine-1-carboxylic acid 2-oxo-, 2-propenyl ester, TFA, triphosgene, NaHCO₃ (sat.), and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a white solid. LC-MS: 501 (M⁺+1).
 5
¹H NMR (CDCl₃) δ 7.70 (d, 1H), 7.60 (s, 1H), 7.15 (d, 1H), 6.0 (s, 2H), 5.40 (d, 1H), 5.30 (m, 2H), 4.80 (m, 5H), 4.40 (m, 1H), 3.65 (m, 4H), 3.40 (m, 1H), 3.10 (m, 4H), 2.10 (m, 1H), 1.95 (m, 3H), 1.60 (m, 1H), 1.50 (m, 1H).

10 **Example 230 :**

Preparation of (3*S*)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-carboxylic acid phenyl ester



15

As described for example 213, (3*R*)-3-[(1,1-dimethylethoxy)carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-carboxylic acid, phenyl ester, TFA, triphosgene, NaHCO₃ (sat.), and 7-chloro-4-(1-piperazinyl)-2-

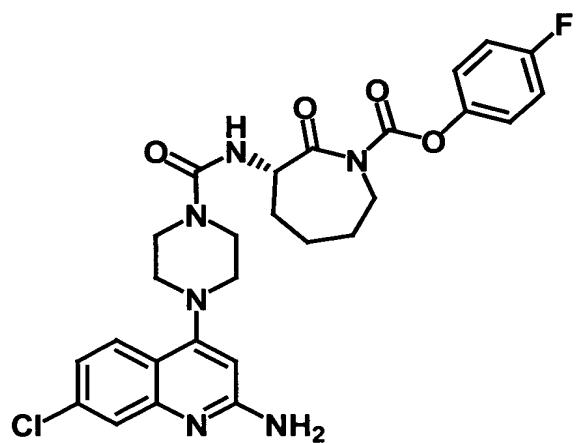
quinolinamine are reacted to afford the product as a light yellow solid. LC-MS: 537 (M^++1). 1H NMR ($CDCl_3$) δ 1.56~2.08 (m, 5H), 2.16~2.24 (m, 1H), 3.14 (m, 4H), 3.54 (dd, 1H), 3.68 (m, 4H), 4.54 (br.dd, 1H), 4.84 (br.s, 2H), 4.90 (ddd, 1H), 5.80 (d, 1H), 6.15 (s, 1H), 7.16~7.22 (m, 3H), 7.27 (m, 1H), 7.41 (m, 2H), 7.62 (d, 1H), 7.72 (d, 1H).

5

Example 231 :

Preparation of (3S)-3-[[(4-(2-Amino-7-chloro-4-quinoliny)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-carboxylic acid-4-fluorophenyl ester

10

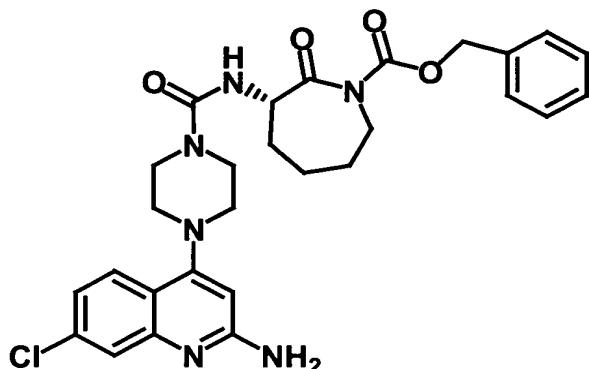


As described for example 213, (3*R*)-3-[[(1,1-dimethylethoxy)carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-carboxylic acid-4-fluorophenyl ester, TFA, triphosgene, $NaHCO_3$ (sat.), and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a light yellow solid. LC-MS: 555 (M^++1). 1H NMR ($CDCl_3$): δ 1.54~2.10 (m, 5H), 2.16~2.24 (m, 1H), 3.14 (m, 4H), 3.54 (dd, 1H), 3.68 (m, 4H), 4.52 (dd, 1H), 4.84 (br.s, 2H), 4.90 (ddd, 1H), 5.94 (d, 1H), 6.15 (s, 1H), 7.06~7.13 (m, 2H), 7.13~7.20 (m, 3H), 7.62 (d, 1H), 7.72 (d, 1H).

15

Example 232 :

Preparation of (3S)-[[[4-(2-Amino-7-chloro-4-quinoliny)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-carboxylic acid, phenylmethyl ester

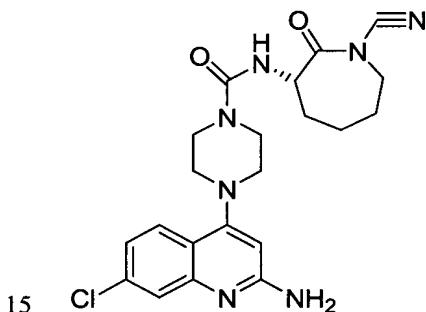


As described example 78, (3R)-3-aminohexahydro-2-oxo-, phenylmethyl ester-1H-azepine-1-carboxylic acid obtained by de-protection of (3S)-3-[(1,1-dimethylethoxy)carbonyl]amino]hexahydro-2-oxo-1H-azepine-1-carboxylic acid,

- 5 phenylmethyl ester with TFA, p-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a white solid. LC-MS: 551 (M^++1). ^1H NMR (CDCl_3) δ 1.50 (m, 1H), 1.65 (m, 1H), 1.92 (m, 3H), 2.13 (d, 1H), 3.10 (m, 4H), 3.40 (m, 1H), 3.65 (m, 4H), 4.45 (m, 1H), 4.85 (m, 3H), 5.30 (s, 2H), 6.00 (d, 1H), 6.15 (s, 1H), 7.20 (d, 1H), 7.40 (m, 5H), 7.60 (s, 1H),
10 7.70 (d, 1H).

Example 233 :

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-1-cyanohexahydro-2-oxo-1H-azepin-3-yl]-1-piperazinecarboxamide

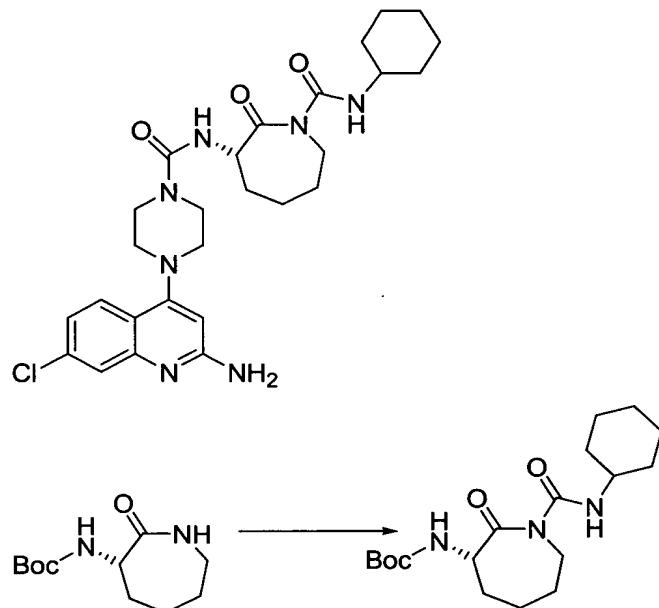


- 15 As described example 78, (3R)-3-amino-1-cyanohexahydro-2H-azepin-2-one obtained by de-protection of [(3S)-1-cyanohexahydro-2-oxo-1H-azepin-3-yl]-carbamic acid-1,1-dimethylethyl ester with TFA, p-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a white solid. LC-MS: 551 (M^++1). ^1H NMR (DMSO-d_6): δ 1.6(m, 2H), 1.92 (m, 3H), 3.23 (br. s,

4H), 3.61 (br. s, 4H), 3.77 (m, 1H), 3.95 (dd, 1H), 4.58 (t, 1H), 6.38 (s, 1H), 7.0 (d, 1H), 7.43 (d, 1H), 7.62 (s, 1H), 7.9 (d, 1H), 8.41 (br., s, 1H).

Example 234 :

5 Preparation of (3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-N-cyclohexylhexahydro-2-oxo-1*H*-azepine-1-carboxamide



10

Preparation of [(3S)-1-[(cyclohexylamino)carbonyl]hexahydro-2-oxo-1*H*-azepin-3-yl]carbamic acid-1,1-dimethylethyl ester (A. D. Borthwick, et al. *J. Med. Chem.*, **2000**, 43, 4452-4464): To a stirred solution of [(3S)-hexahydro-2-oxo-1*H*-azepin-3-yl]-carbamic acid-1,1-dimethylethyl ester (500 mg, 2.2 mmol) in THF (10 mL) at -78 °C was added 15 LiHMDS (4.4 ml, 4.4 mmol, 1.0 M in THF). After addition, dry-ice bath was removed, and the reaction was stirred at room temperature for about 30 min. The reaction was re-cooled to -78 °C, and cyclohexyl isocyanate (554 mg, 4.4 mmol) was added. The reaction was allowed to warm to rt over 2 h, and was quenched with NH₄Cl (sat., 10 mL). The reaction mixture was extracted with EtOAc (3x30 mL), washed with brine (15 mL), and dried over Na₂SO₄. Concentration *in vacuo* followed by purification through flash chromatography (hexane:EtOAc/ 5:1) afforded the title product. ¹H NMR

20

(CDCl₃) δ 1.1-1.4 (m, 6H), 1.45 (s, 9H), 1.5-1.7 (m, 3H), 1.75 (m, 1H), 1.96 (m, 2H), 2.28 (dq, 1H), 3.46 (br. s, 1H), 4.41 (br. s, 1H), 4.51 (tt, 1H).

In a similar manner following intermediates were prepared:

5

[(3*R*)-1-(aminocarbonyl)hexahydro-2-oxo-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester

[(3*R*)-hexahydro-2-oxo-1-[(2,2,2-trifluoroethyl)amino]carbonyl]-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester

10 [(3*R*)-1-[(cyclopropylamino)carbonyl]hexahydro-2-oxo-1*H*-azepin-3-yl]-carbamic acid-1,1-dimethylethyl ester

[(3*R*)-hexahydro-2-oxo-1-[(propylamino)carbonyl]-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester

(3*S*)-hexahydro-2-oxo-1-(1-pyrrolidinylcarbonyl)-1*H*-azepin-3-yl]-carbamic acid,

15 1,1-dimethylethyl ester

[(3*S*)-1-[(cyclopentylamino)carbonyl]hexahydro-2-oxo-1*H*-azepin-3-yl]- carbamic acid-1,1-dimethylethyl ester. ¹H NMR (CDCl₃) δ 1.36 (m, 2H), 1.41 (s, 9H), 1.5-1.7 (m, 5H), 1.87 (m, 4H), 2.12 (m, 2H), 3.86 (m, 1H), 4.44 (br. s, 1H), 5.17 (m, 1H).

[(3*S*)-1-[(cyclobutylamino)carbonyl]hexahydro-2-oxo-1*H*-azepin-3-yl] carbamic

20 acid-1,1-dimethylethyl ester. ¹H NMR (CDCl₃) δ 1.4 (m, 1H), 1.43 (s, 9H), 1.5-1.98 (m, 6 H), 2.1 (m, 1H), 2.38 (m, 2H), 3.17 (m, 1H), 4.33 (m, 1H), 4.64 (m, 1H), 4.81 (m, 1H), 5.61 (br. s, 1H), 9.25 (br. s, 1H).

[(3*S*)-1-[(4-fluorophenyl)amino]carbonyl]-hexahydro-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester

25 [(3*S*)-1-[(ethylamino)carbonyl]hexahydro-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester

[(3*S*)-hexahydro-2-oxo-1-[(phenylamino)carbonyl]-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester

30 [(3*S*)-hexahydro-1-[(1-methylethyl)amino]carbonyl]-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester

[(3*S*)-1-[(butylamino)carbonyl]hexahydro-2-oxo-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester

[(3S)-hexahydro-1-[[[(2-methylpropyl)amino]carbonyl]-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester

[(3S)-1-[[[(1,1-dimethylethyl)amino]carbonyl]hexahydro-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester

5 [(3S)-1-[(dimethylamino)carbonyl]hexahydro-2-oxo-1*H*-azepin-3-yl]- carbamic acid-1,1-dimethylethyl ester

(3S)-hexahydro-2-oxo-1-[[[4-(trifluoromethyl)phenyl]amino]carbonyl]-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester

10 [(3S)-1-[[[(3-fluorophenyl)amino]carbonyl]hexahydro-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester

[(3S)-hexahydro-1-[[[(4-methylphenyl)amino]carbonyl]-2-oxo-1*H*-azepin-3-yl]- carbamic acid 1,1-dimethylethyl ester

[(3S)-1-[[[(2-fluorophenyl)amino]carbonyl]hexahydro-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester

15 [(3S)-1-[[[(2,6-diethylphenyl)amino]carbonyl]hexahydro-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester

[(3S)-hexahydro-1-[(methylamino)carbonothioyl]-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester

20 [(3S)-1-[(ethylamino)carbonothioyl]hexahydro-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester

[(3S)-hexahydro-1-[[[(1-methylethyl)amino]carbonothioyl]-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester

[(3S)-hexahydro-2-oxo-1-[(2-propenylamino)carbonothioyl]-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester

25

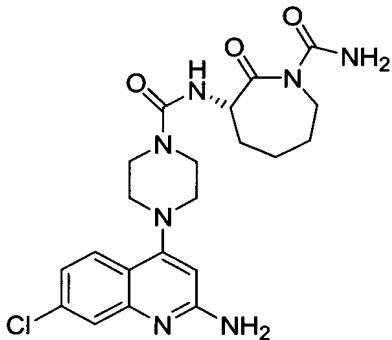
*Preparation of (3S)-3-[[[4-(2-amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-N-cyclohexylhexahydro-2-oxo-1*H*-azepine-1-carboxamide:*
As described example 78, (3*R*)-3-amino-N-cyclohexylhexahydro-2-oxo-1*H*-azepine-1-carboxamide obtained by de-protection of [(3S)-1-

30 [(cyclohexylamino)carbonyl]hexahydro-2-oxo-1*H*-azepin-3-yl]-carbamic acid-1,1-dimethylethyl ester with TFA, *p*-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a white solid. ¹H NMR (DMSO-d₆): δ 1.2 (m, 4H), 1.5 (m, 1H), 1.6-1.8 (m, 4H), 2.97 (m, 4H),

3.4(m, 1H), 3.48 (m, 4H), 6.21 (s, 1H), 6.27 (d, 1H), 6.43 (m, 1H), 7.11 (d, 1H), 7.37 (s, 1H), 7.73 (d, 1H).

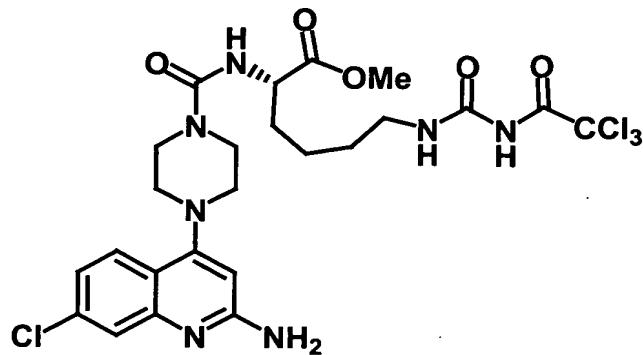
Example 235 :

5 Preparation of (3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-carboxamide



10 Preparation of (3S)-3-[[[4-(2-amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-1*H*-azepine-1-carboxamide:
As described for example 213, [(3S)-1-(aminocarbonyl)hexahydro-2-oxo-1*H*-azepin-3-yl]-carbamic acid-1,1-dimethylethyl ester, TFA, triphosgene, NaHCO₃ (sat.), and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a white solid. LC-MS: 460 (M⁺+1). ¹H NMR (DMSO-d₆+TFA): δ 1.16~1.30 (m, 1H), 1.54~1.88 (m, 5H), 3.26 (m, 4H), 3.60 (m, 4H), 4.56 (ddd, 1H), 4.70 (br.dd, 1H), 6.32 (s, 1H), 7.38~7.46 (m, 2H), 7.62 (d, 1H), 7.88 (d, 1H), 8.20~8.50 (m, 3H).

15 (2S)-2-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-6-[[[(trichloroacetyl)amino]carbonyl]amino]-hexanoic acid, methyl ester



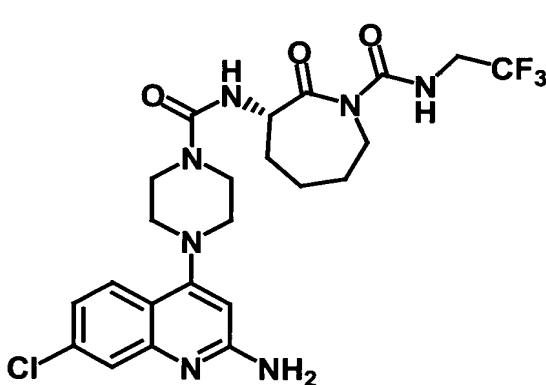
The title product was obtained as a product in example 235. LC-MS: 636 (M^++1).

^1H NMR (CDCl_3): δ 1.38~1.98 (m, 6H), 3.14 (m, 4H), 3.37 (dd, 2H), 3.68 (m, 4H), 3.78 (s, 3H), 4.56 (m, 1H), 4.98 (br.s, 1H), 5.14 (d, 1H), 6.16 (s, 1H), 7.18 (dd, 1H), 7.26 (s,

5 1H), 7.64 (d, 1H), 7.72 (d, 1H), 7.84 (m, 1H).

Example 236 :

Preparation of (3S)-3-[[[4-(2-Amino-7-chloro-4-quinoliny)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-N-(2,2,2-trifluoroethyl)-1*H*-azepine-10 1-carboxamide

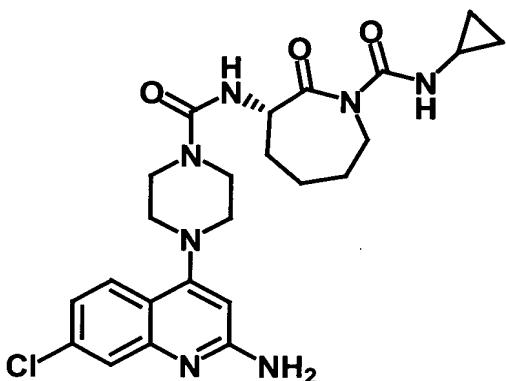


As described for example 213, [(3*R*)-hexahydro-2-oxo-1-[(2,2,2-trifluoroethyl)amino]carbonyl]-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester,

15 TFA, triphosgene, NaHCO_3 (sat.), and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a light yellow solid. LC-MS: 542 (M^++1). ^1H NMR (CDCl_3): δ 1.44 (m, 1H), 1.64 (m, 1H), 1.82~2.02 (m, 4H), 3.14 (m, 4H), 3.31(dd, 1H), 3.68 (m, 4H), 3.98 (m, 2H), 4.84 (dd, 1H), 4.88~5.00 (m, 3H), 5.80 (d, 1H), 6.03 (s, 1H), 7.16 (dd, 1H), 7.60 (d, 1H), 7.71 (d, 1H), 9.54 (t, 1H).

Example 237 :

Preparation of (3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-N-cyclopropylhexahydro-2-oxo-1*H*-azepine-1-carboxamide



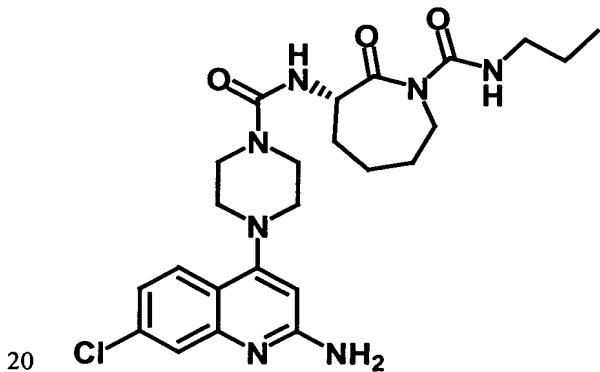
As described for example 213, [(3*R*)-1-[(cyclopropylamino)carbonyl]hexahydro-2-oxo-1*H*-azepin-3-yl]-carbamic acid-1,1-dimethylethyl ester, TFA, triphosgene, NaHCO₃ (sat.), and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product

as a white solid. LC-MS: 500 (M⁺+1). ¹H NMR (CDCl₃): δ 0.50~0.62 (m, 2H), 0.72~0.84 (m, 2H), 1.42 (m, 1H), 1.60 (m, 1H), 1.82~2.02 (m, 3H), 2.13 (m, 1H), 2.76 (ddd, 1H), 3.12 (br.dd, 4H), 3.24 (br.dd, 1H), 3.66 (m, 4H), 4.82~4.94 (m, 2H), 5.02 (br.s, 2H), 5.86 (d, 1H), 6.16 (s, 1H), 7.16 (dd, 1H), 7.60 (d, 1H), 7.72 (d, 1H), 9.10 (d, 1H).

15

Example 238 :

Preparation of (3S)-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-N-propyl-1*H*-azepine-1-carboxamide

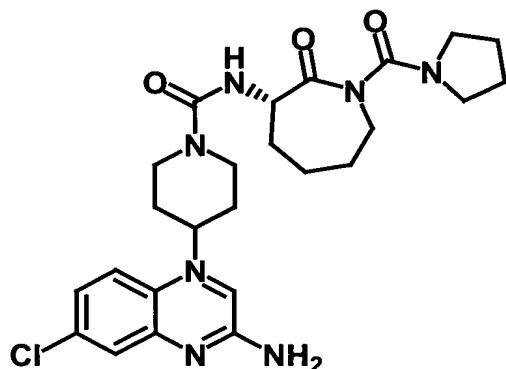


As described for example 213, [(3*R*)-hexahydro-2-oxo-1-[(propylamino)carbonyl]-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester, TFA, triphosgene, NaHCO₃ (sat.), and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a white solid. LC-MS: 502 (M⁺+1). ¹H NMR (CD₃OD) δ 0.95 (t, 3H), 1.55 (m, 4H), 1.90 (m, 3H), 2.10 (d, 1H), 3.10 (m, 4H), 3.25 (m, 3H), 3.65 (m, 4H), 4.80-4.95 (m, 4H), 5.86 (d, 1H), 6.12 (s, 1H), 7.12 (d, 1H), 7.58 (s, 1H), 7.65 (d, 1H), 9.00 (m, 1H).

10

Example 239 :

Preparation of 4-(2-Amino-7-chloro-4-quinoliny)-N-[(3*S*)-hexahydro-2-oxo-1-(1-pyrrolidinylcarbonyl)-1*H*-azepin-3-yl]-1-piperazinecarboxamide



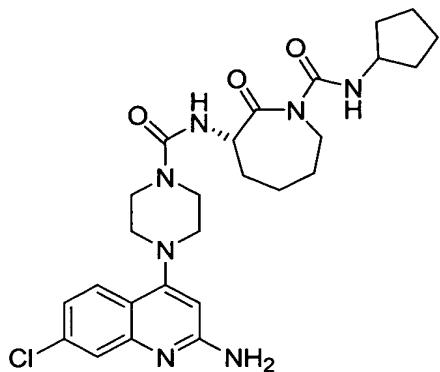
15

As described for example 78, (3*R*)-3-aminohexahydro-1-(1-pyrrolidinylcarbonyl)-2*H*-azepin-2-one obtained by de-protection of (3*S*)-hexahydro-2-oxo-1-(1-pyrrolidinylcarbonyl)-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester with TFA, diisopropyl(ethyl)amine, and 7-chloro-4-(piperazin-1-yl)quinoline are reacted to give the product. LC-MS: 514 (M⁺+1). ¹H NMR (CDCl₃) δ 7.70 (d, 1H), 7.62 (s, 1H), 7.18 (d, 1H), 6.15 (s, 1H), 6.00 (s, 1H), 5.00 (s, 2H), 4.60 (m, 1H), 3.75 (m, 1H), 3.65 (m, 4H), 3.50 (m, 3H), 3.35 (m, 2H), 3.15 (m, 4H), 1.80-2.20 (m, 7H), 1.60 (m, 2H), 1.30 (m, 1H).

25 **Example 240 :**

192

Preparation of (3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-N-cyclopentylhexahydro-2-oxo-1*H*-azepine-1-carboxamide

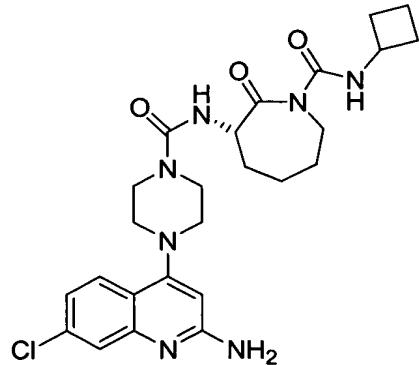


5 As described example 78, (3R)-3-amino-N-cyclopentylhexahydro-2-oxo-1*H*-azepine-1-carboxamide obtained by de-protection of [(3S)-1-[(cyclopentylamino)carbonyl]hexahydro-2-oxo-1*H*-azepin-3-yl]-carbamic acid-1,1-dimethylethyl ester with TFA, p-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a white solid. ¹H NMR (DMSO-d₆): δ 1.38 (m, 2H), 1.63 (m, 5H), 2.06 (m, 2H), 3.15 (m, 4), 3.6 (m, 4H), 4.16 (m, 1H), 4.41 (d, 1H), 4.78 (br. s, 2H), 6.18 (s, 1H), 7.2 (d, 1H), 7.62 (s, 1H), 7.73 (d, 1H).

10

Example 241 :

15 **Preparation of (3S)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-N-cyclobutylhexahydro-2-oxo-1*H*-azepine-1-carboxamide**



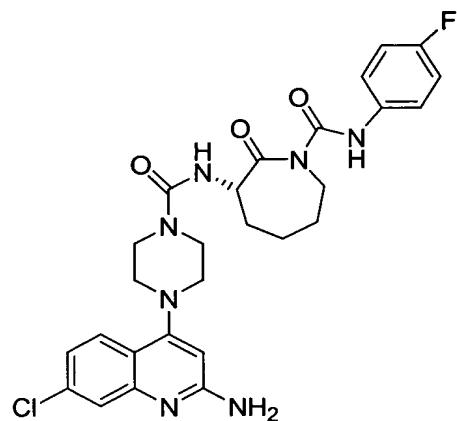
As described example 78, (3R)-3-amino-N-cyclobutylhexahydro-2-oxo-1*H*-azepine-1-carboxamide by de-protection of [(3S)-1-[(cyclobutylamino)carbonyl]hexahydro-2-oxo-

20

1*H*-azepin-3-yl] carbamic acid-1,1-dimethylethyl ester with TFA, p-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product. ^1H NMR (DMSO- d_6): δ 1.24 (br. s, 1*H*), 1.5-1.9 (m, 7*H*), 2.2 (m, 2*H*), 2.99 (m, 4*H*), 3.6 (m, 4*H*), 4.17 (m, 1*H*), 4.52 (m, 1*H*), 4.66 (t, 1*H*), 5 6.14 (s, 1*H*, 6.42 (br. s, 1*H*), 6.8 (d, 1*H*), 7.1 (d, 1*H*), 7.39 (s, 1*H*), 7.76 (d, 1*H*), 9.3 (d, 1*H*).

Example 242 :

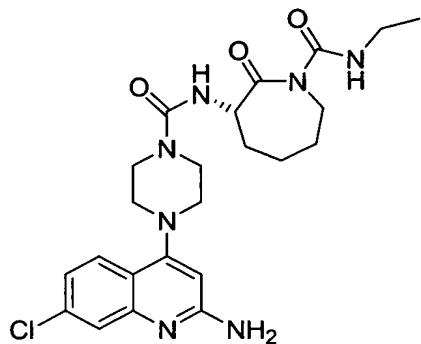
Preparation of (*3R*)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-
10 piperazinyl]carbonyl]amino]-*N*-(4-fluorophenyl)hexahydro-2-oxo-1*H*-azepine-1-
carboxamide



As described for example 78, (*3R*)-3-amino-*N*-(4-fluorophenyl)hexahydro-2-oxo-1*H*-azepine-1-carboxamide obtained by de-protection of [(*3S*)-1-[[4-(
15 fluorophenyl)amino]carbonyl]hexahydro-2-oxo-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester with TFA, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to give the product. LC-MS: 554 (M^++1). ^1H NMR (CDCl_3) δ 1.4-2.25 (m, 6*H*), 3.25-3.45 (m, 4*H*), 3.7-3.9 (m, 4*H*), 4.85-5.1 (m, 2*H*), 5.85-6.0 (s, 20 1*H*), 6.1-6.2 (m, 1*H*), 7.0-7.08 (m, 2*H*), 7.32-7.38 (m, 1*H*), 7.44-7.5 (m, 2*H*), 7.68-7.76 (m, 2*H*), 11.2(s, 1*H*).

Example 243 :

Preparation of (*3S*)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-
25 piperazinyl]carbonyl]amino]-*N*-ethylhexahydro-2-oxo-1*H*-azepine-1-carboxamide

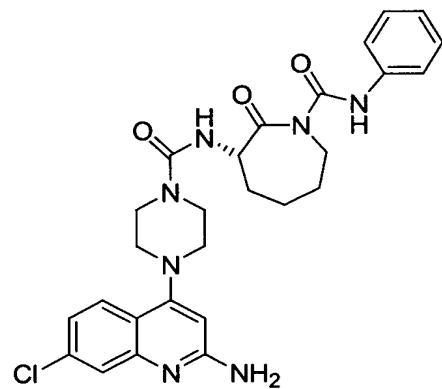


As described for example 78, (3*R*)-3-amino-*N*-ethylhexahydro-2-oxo-1*H*-azepine-1-carboxamide obtained by the de-protection of [(3*S*)-1-

- 5 [(ethylamino)carbonyl]hexahydro-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester with TFA, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to give the product. LC-MS: 487 (M^++1). ^1H NMR (CDCl_3) δ 1.2 (t, 3H), 1.3-1.7 (m, 2H), 1.7-2.2 (m, 4H), 3.15-3.45 (m, 7H), 3.6-3.8 (m, 4H), 4.8-5.0 (m, 2H), 5.8-6.0 (m, 1H), 6.0-6.2 (s, 1H), 7.34-7.4 (m, 1H), 7.7 (d, 1H), 7.8 (m, 1H),
10 8.9-9.0 (m, 1H).

Example 244 :

- 15 Preparation of (3*S*)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-*N*-phenyl-1*H*-azepine-1-carboxamide

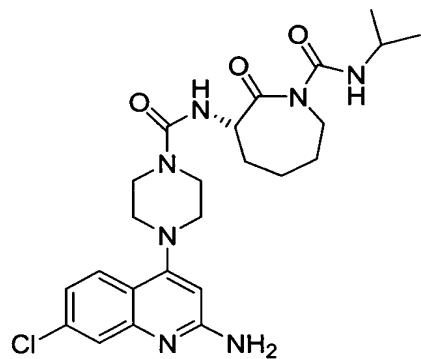


- 20 As described for example 78, (3*R*)-3-aminohexahydro-2-oxo-*N*-phenyl-1*H*-azepine-1-carboxamide obtained by de-protection of [(3*S*)-hexahydro-2-oxo-1-[(phenylamino)carbonyl]-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester with

TFA, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to give the product. LC-MS: 535 (M^++1). 1H NMR ($CDCl_3$): δ 1.3-1.8 (m, 2H), 1.8-2.3 (m, 4H), 3.1-3.25 (m, 4H), 3.26-3.34 (m, 1H), 3.6-3.8 (m, 4H), 4.76-4.86 (s, 2H), 4.88-5.06 (m, 2H), 5.8-6.1 (m, 1H), 6.1-6.2 (s, 1H), 7.09-7.15 (m, 1H), 7.16-7.25 (m, 1H), 7.3-7.37 (m, 2H), 7.5-7.55 (m, 1H), 7.62-7.64 (d, 1H), 7.72-7.76 (d, 1H), 11.2-11.4 (s, 1H).

Example 245 :

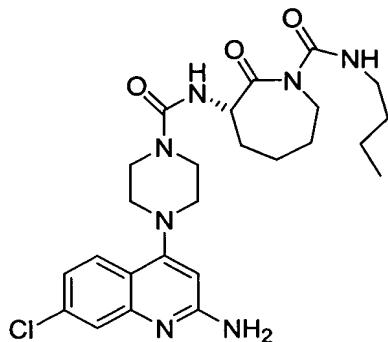
Preparation of (3*S*)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-N-(1-methylethyl)-2-oxo-1*H*-azepine-1-carboxamide



As described for example 78, (3*R*)-3-aminohexahydro-N-(1-methylethyl)-2-oxo-1*H*-azepine-1-carboxamide obtained by de-protection of [(3*S*)-hexahydro-1-[(1-methylethyl)amino]carbonyl]-2-oxo-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester with TFA, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to give the product. LC-MS: 501 (M^++1). 1H NMR ($CDCl_3$): δ 1.2 (m, 6H), 1.3-1.5 (m, 1H), 1.5-1.7 (m, 1H), 1.8-2.2 (m, 4H), 3.1-3.3 (m, 5H), 3.6-3.8 (m, 4H), 3.9-4.1 (m, 1H), 4.78-5.0 (m, 4H), 5.8-5.9 (m, 1H), 6.1-6.2 (s, 1H), 7.1-7.2 (m, 1H), 7.6-7.64 (d, 1H), 7.7-7.76 (m, 1H), 8.9-8.96 (d, 1H).

Example 246 :

Preparation of (3*S*)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-*N*-butylhexahydro-2-oxo-1*H*-azepin-1-carboxamide

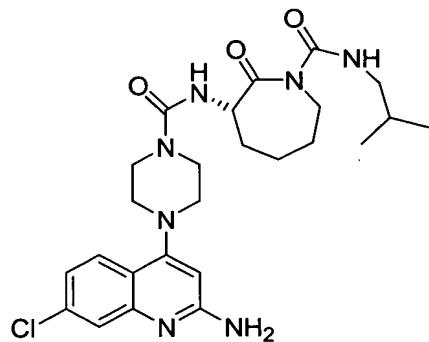


As described for example 78, (3*R*)-3-amino-*N*-butylhexahydro-2-oxo-1*H*-azepine-1-carboxamide obtained by de-protection of [(3*S*)-1-[(butylamino)carbonyl]hexahydro-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester with TFA,

5 diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to give the product. ^1H NMR (CDCl_3): δ 0.9-1.0 (t, 3H), 1.3-2.4 (m, 10H), 3.1-3.2 (m, 4H), 3.2-3.4 (m, 3 H), 3.6-3.76 (m, 4H), 4.78-5.0 (m, 4H), 5.8-5.9 (m, 1H), 6.1-6.2 (s, 1H), 7.14-7.2 (m, 1H), 7.6-7.64 (d, 1H), 7.7-7.8 (d, 1H), 8.98-9.06 (m, 1H).

10 **Example 247 :**

Preparation of (3*S*)-3-[[[4-(2-Amino-7-chloro-4-quinoliny)-1-piperazinyl]carbonyl]amino]hexahydro-*N*-(2-methylpropyl)-2-oxo-1*H*-azepine-1-carboxamide

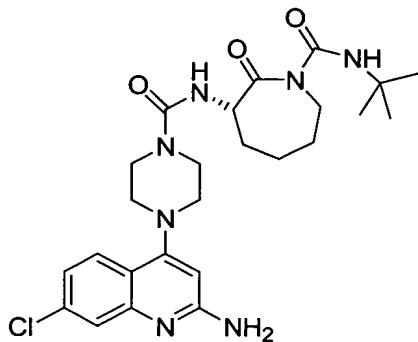


15 As described for example 78, (3*R*)-3-aminohexahydro-*N*-(2-methylpropyl)-2-oxo-1*H*-azepine-1-carboxamide obtained by de-protection of [(3*S*)-hexahydro-1-[(2-methylpropyl)amino]carbonyl]-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester with TFA, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to give the product. LC-MS: 515 (M^++1). ^1H NMR (CDCl_3): δ 0.9-1.1(d, 6H), 1.3-2.3 (m, 7H), 3.1-3.3 (m, 3H), 3.3-3.4 (m, 4 H), 3.64-3.8 (m, 4H), 4.78-5.0 (m, 20

2H), 5.76-5.84 (m, 1H), 6.1-6.2 (s, 1H), 7.3-7.4 (m, 1H), 7.66-7.74 (d, 1H), 7.75-7.8 (d, 1H), 9.04-9.14 (m, 1H).

5 **Example 248 :**

Preparation of (3*S*)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-*N*-(1,1-dimethylethyl)hexahydro-2-oxo-1*H*-azepine-1-carboxamide

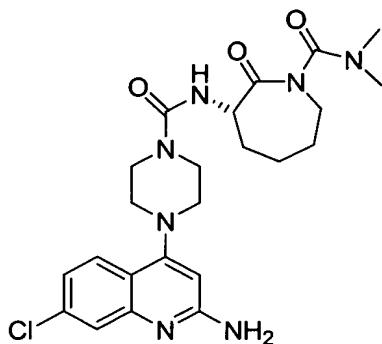


10 As described for example 78, (3*R*)-3-amino-*N*-(1,1-dimethylethyl)hexahydro-2-oxo-1*H*-azepine-1-carboxamide obtained by the de-protection of [(3*S*)-1-[(1,1-dimethylethyl)amino]carbonyl]hexahydro-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester with TFA, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to give the product. LC-MS: 515 ($M^+ + 1$). ^1H NMR (CDCl_3): δ 1.3-1.5 (s, 9H), 1.5-2.3 (m, 6H), 3.1-3.29 (m, 5 H), 3.6-3.75 (m, 4H), 4.76-5.0 (m, 4H), 5.8-6.0 (m, 1H), 6.1-6.2 (s, 1H), 7.15-7.2 (m, 1H), 7.6-7.64 (d, 1H), 7.70-7.75 (d, 1H), 9.0-9.2 (s, 1H).

15

Example 249 :

20 **Preparation of (3*S*)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-*N,N*-dimethyl-2-oxo-1*H*-azepine-1-carboxamide**



As described for example 78, (3*R*)-3-aminohexahydro-*N,N*-dimethyl-2-oxo-1*H*-azepine-1-carboxamide obtained by de-protection of [(3*S*)-1-

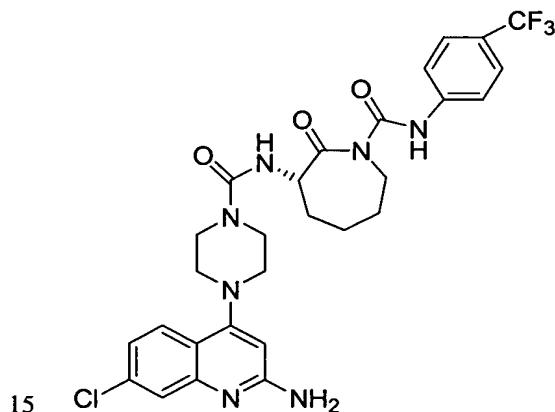
[(dimethylamino)carbonyl]hexahydro-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-

5 dimethylethyl ester with TFA, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to give the product. LC-MS: 587 ($M^{+}+1$). 1H NMR ($CDCl_3$): δ 1.36-2.2 (m, 5H), 2.6-3.2 (m, 8H), 3.22-3.4 (m, 4H), 3.4-3.9 (m, 4H), 4.4-4.8 (m, 2H), 5.9-6.2 (m, 2H), 6.4-6.6 (m, 1H), 7.3-7.4 (m, 1H), 7.65-7.73 (m, 1H), 7.74-7.8 (m, 1H).

10

Example 250 :

Preparation of (3*S*)-3-[[[4-(2-Amino-7-chloro-4-quinoliny)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-*N*-[4-(trifluoromethyl)phenyl]- 1*H*-azepine-1-carboxamide



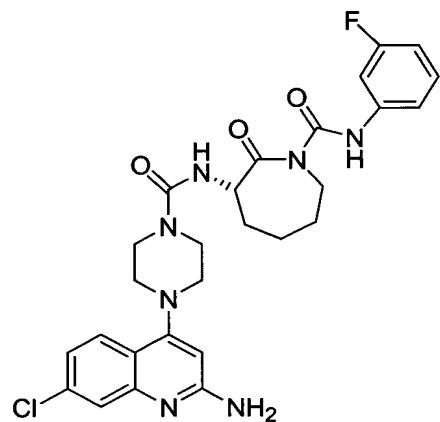
15

As described for example 78, (3*R*)-3-aminohexahydro-2-oxo-*N*-[4-(trifluoromethyl)phenyl]- 1*H*-azepine-1-carboxamide obtained by de-protection of [(3*S*)-hexahydro-2-oxo-1-[[[4-(trifluoromethyl)phenyl]amino]carbonyl]-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester with TFA, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to give the product. LC-MS: 603 ($M^{+}+1$).

¹H NMR (CDCl₃): δ 1.3-1.8 (m, 2H), 1.8-2.3 (m, 4H), 3.1-3.25 (m, 4H), 3.26-3.4 (m, 1H), 3.6-3.8 (m, 4H), 4.76-5.1 (m, 4H), 5.7-5.9 (m, 1H), 6.1-6.2 (s, 1H), 7.16-7.2 (m, 1H), 7.56-7.68 (m, 5H), 7.7-7.76 (d, 1H), 11.4-11.6 (s, 1H).

5 **Example 251 :**

Preparation of (3*S*)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-*N*-(3-fluorophenyl)hexahydro-2-oxo-1*H*-azepine-1-carboxamide

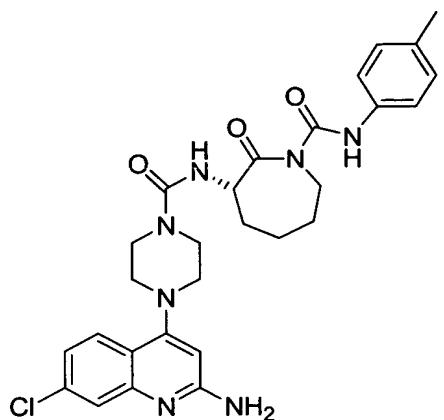


10 As described for example 78, (3*R*)-3-amino-*N*-(3-fluorophenyl)hexahydro-2-oxo-1*H*-azepine-1-carboxamide obtained by de-protection of [(3*S*)-1-[[3-fluorophenyl]amino]carbonyl]hexahydro-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester with TFA, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to give the product. LC-MS: 553 (M⁺+1). ¹H NMR (CDCl₃): δ 1.3-1.8 (m, 2H), 1.8-2.3 (m, 4H), 3.1-3.25 (m, 4H), 3.26-3.4 (m, 1H), 3.6-3.8 (m, 4H), 4.8-5.1 (m, 4H), 5.8-5.9 (m, 1H), 6.1-6.2 (s, 1H), 6.78-6.86 (m, 1H), 7.12-7.2 (m, 2H), 7.22-7.31 (m, 1H), 7.43-7.5 (m, 1H), 7.6-7.64 (d, 1H), 7.7-7.76 (d, 1H), 11.3-11.5 (s, 1H).

15

20 **Example 252 :**

Preparation of (3*S*)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-*N*-(4-methylphenyl)-2-oxo-1*H*-azepine-1-carboxamide



As described for example 78, (3*R*)-3-aminohexahydro-*N*-(4-methylphenyl)-2-oxo-1*H*-azepine-1-carboxamide obtained by de-protection of [(3*S*)-hexahydro-1-[(4-methylphenyl)amino]carbonyl]-2-oxo-1*H*-azepin-3-yl]-carbamic acid 1,1-dimethylethyl ester with TFA, diisopropyl(ethyl)amine, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to give the product. LC-MS: 549 (M^++1). 1H NMR ($CDCl_3$): δ 1.3-2.23 (m, 6H), 2.31-2.4 (s, 3H), 3.1-3.22 (m, 4H), 3.28-3.4 (m, 1H), 3.6-3.8 (m, 4H), 4.8-5.1 (m, 4H), 5.8-5.9 (m, 1H), 6.1-6.2 (s, 1H), 7.1-7.2 (m, 3H), 7.37-7.43 (m, 2H), 7.61-7.64 (d, 1H), 7.7-7.76 (d, 1H), 11.1-11.3 (s, 1H).

10

Example 253 :

Preparation of (3*S*)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]-*N*-(2-fluorophenyl)hexahydro-2-oxo-1*H*-azepine-1-carboxamide

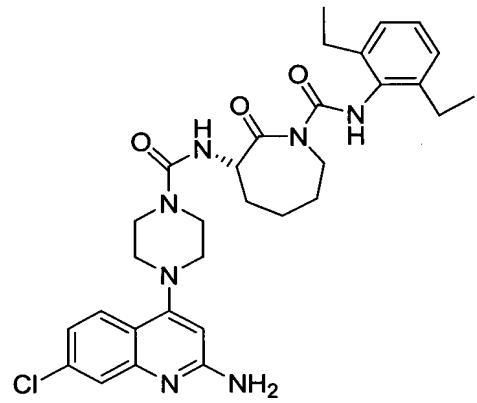
15 As described for example 78, (3*R*)-3-amino-*N*-(2-fluorophenyl)hexahydro-2-oxo-1*H*-azepine-1-carboxamide obtained by de-protection of [(3*S*)-1-[(2-fluorophenyl)amino]carbonyl]hexahydro-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester with TFA, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-

quinolinamine are reacted to give the product. LC-MS: 553 (M^++1). 1H NMR ($CDCl_3$): δ 1.2-2.3 (m, 6H), 3.1-3.24 (m, 4H), 3.28-3.44 (m, 1H), 3.6-3.84 (m, 4H), 4.7-4.88 (s, 2H), 4.9-5.1 (m, 2H), 5.8-5.9 (m, 1H), 6.16-6.24 (s, 1H), 7.02-7.24 (m, 4H), 7.61-7.65 (d, 1H), 7.71-7.78 (d, 1H), 8.2-8.28 (m, 1H), 11.5-11.7 (s, 1H).

5

Example 254 :

Preparation of (3S)-3-[[[4-(2-Amino-7-chloro-4-quinoliny)-1-piperazinyl]carbonyl]amino]-N-(2,6-diethylphenyl)hexahydro-2-oxo-1*H*-azepine-1-carboxamide



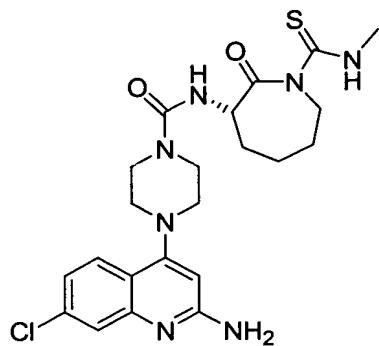
10

As described for example 78, (3*R*)-3-amino-N-(2,6-diethylphenyl)hexahydro-2-oxo-1*H*-azepine-1-carboxamide obtained by de-protection of [(3*S*)-1-[(2,6-diethylphenyl)amino]carbonyl]hexahydro-2-oxo-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester with TFA, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to give the product. LC-MS: 591 (M^++1). 1H NMR ($CDCl_3$): δ 1.2 (t, 6H), 1.3-2.3 (m, 6H), 2.6 (q, 2H), 3.1-3.23 (m, 4H), 3.3-3.4 (m, 1H), 3.6-3.8 (m, 4H), 4.8-5.1 (m, 4H), 5.8-6.0 (m, 1H), 6.1-6.2 (s, 1H), 7.1-7.28 (m, 4H), 7.6-7.64 (d, 1H), 7.70-7.75 (d, 1H), 10.36-10.48 (s, 1H).

20

Example 255 :

Preparation of 4-(2-Amino-7-chloro-4-quinoliny)-N-[(3*S*)-hexahydro-1-[(methylamino)carbonothioyl]-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide:



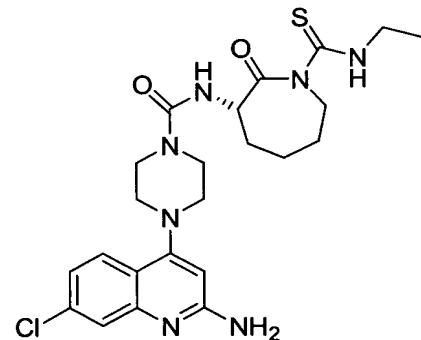
As described for example 78, obtained (3*R*)-3-aminohexahydro-*N*-methyl-2-oxo-1*H*-azepine-1-carbothioamide by de-protection of [(3*S*)-hexahydro-1-[(ethylamino)carbonothioyl]-2-oxo-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester with TFA, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to give the product. LC-MS: 489 (M^++1). ^1H NMR (CDCl_3): δ 1.5-2.2 (m, 6H), 3.1-3.17 (m, 4H), 3.17-3.22 (d, 3H), 3.46-3.58 (m, 1H), 3.6-3.74 (m, 4H), 4.8-4.9 (s, 2H), 5.0 (m, 1H), 5.6-5.8 (m, 2H), 6.1-6.2 (s, 1H), 7.14-7.2 (m, 1H), 7.6-7.63 (d, 1H), 7.7-7.74 (d, 1H), 11.2 (m, 1H).

10

Example 256 :

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-*N*-[(3*S*)-1-[(ethylamino)carbonothioyl]hexahydro-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide

15



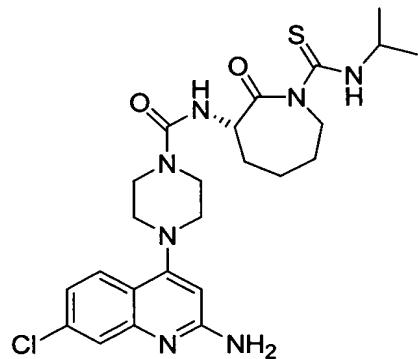
20

As described for example 78, (3*R*)-3-amino-*N*-ethylhexahydro-2-oxo-1*H*-azepine-1-carbothioamide obtained by de-protection of [(3*S*)-1-[(ethylamino)carbonothioyl]hexahydro-2-oxo-1*H*-azepin-3-yl]- carbamic acid, 1,1-dimethylethyl ester with TFA, diisopropyl(ethyl)amine, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to give the product. LC-MS: 503 (M^++1). ^1H NMR (CDCl_3): δ 1.2-1.4 (t, 3H), 1.5-1.78 (m, 2H), 1.8-2.2 (m, 4H), 3.12-

3.18 (m, 4H), 3.44-3.56 (m, 1H), 3.6-3.76 (m, 6H), 4.8-4.9 (s, 2H), 5.0 (m, 1H), 5.64-5.8 (m, 2H), 6.1-6.2 (s, 1H), 7.14-7.2 (m, 1H), 7.6-7.64 (d, 1H), 7.7-7.76 (d, 1H), 11.1-11.3 (m, 1H).

5 **Example 257 :**

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3S)-hexahydro-1-[(1-methylethyl)amino]carbonothioyl]-2-oxo-1*H*-azepin-3-yl]-1-piperazinecarboxamide:



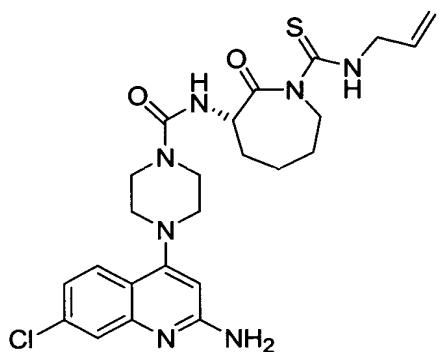
10 As described for example 78, (3*R*)-3-aminohexahydro-*N*-(1-methylethyl)-2-oxo-1*H*-azepine-1-carbothioamide obtained by de-protection of [(3*S*)-hexahydro-1-[(1-methylethyl)amino]carbonothioyl]-2-oxo-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester with TFA, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to give the product. LC-MS: 517 (M^++1). 1H NMR ($CDCl_3$): δ 1.3 (t, 6H), 1.5-2.2 (m, 6H), 3.1-3.17 (m, 4 H), 3.4-3.53 (m, 1H), 3.6-3.75 (m, 4H), 4.49-4.6 (m, 1H), 4.88-4.97 (s, 2H), 4.97-5.05 (m, 1H), 5.6-5.7 (m, 1H), 5.74-5.82 (d, 1H), 6.1-6.2 (s, 1H), 7.14-7.2 (m, 1H), 7.6-7.63 (d, 1H), 7.69-7.74 (d, 1H), 11.0-11.2 (m, 1H).

15

20

Example 258 :

Preparation of 4-(2-Amino-7-chloro-4-quinolinyl)-N-[(3*S*)-hexahydro-2-oxo-1-[(2-propenylamino)carbonothioyl]-1*H*-azepin-3-yl]-1-piperazinecarboxamide



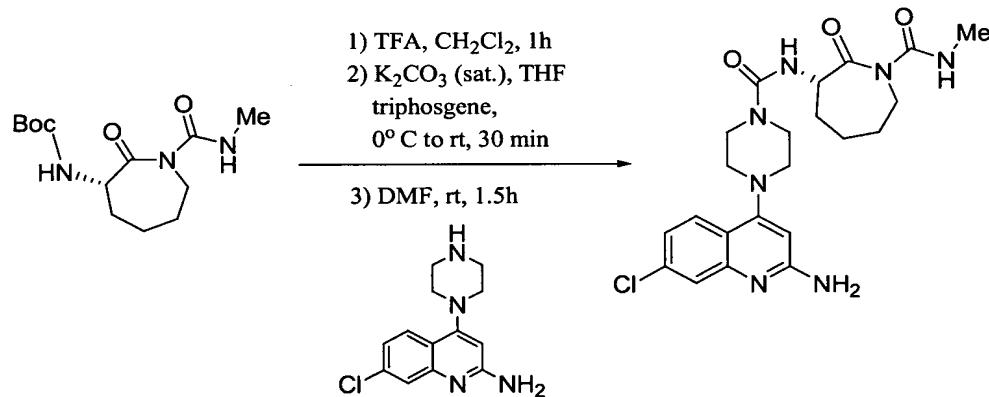
As described for example 78, (3*R*)-3-aminohexahydro-2-oxo-*N*-(2-propenyl)-1*H*-azepine-1-carbothioamide obtained by de-protection of [(3*S*)-hexahydro-2-oxo-1-[(2-propenylamino)carbonothioyl]-1*H*-azepin-3-yl]- carbamic acid,, 1,1-dimethylethyl ester

5 with TFA, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to give the product. LC-MS: 515 (M^++1). ^1H NMR (CDCl_3): δ 1.5-2.3 (m, 6H), 3.1-3.19 (m, 4 H), 3.48-3.58 (m, 1H), 3.6-3.75 (m, 4H), 4.26-4.34 (m, 2H), 4.8-4.9 (s, 2H), 4.99-5.08 (m, 1H), 5.2-5.34 (m, 2H), 5.65-5.78 (m, 2H), 5.9-6.02 (m, 1H), 6.1-6.2 (s, 1H), 7.14-7.2 (m, 1H), 7.6-7.64 (d, 1H), 7.70-7.75 (d, 1H), 11.2-11.4 (m, 1H).

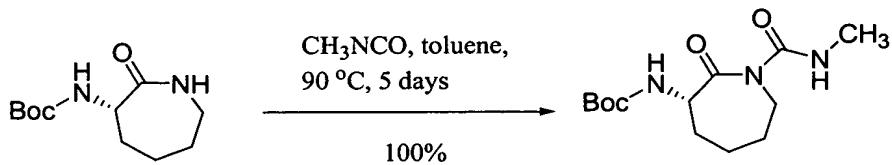
10

Example 259 :

Preparation of (3*S*)-3-[[[4-(2-Amino-7-chloro-4-quinolinyl)-1-piperazinyl]carbonyl]amino]hexahydro-*N*-methyl-2-oxo-1*H*-azepine-1-carboxamide



15



Preparation of [(3R)-Hexahydro-1-[(methylamino)carbonyl]-2-oxo-1H-azepin-3-yl]carbamic acid- 1,1-dimethylethyl ester: A mixture of [(3R)-hexahydro-2-oxo-1H-azepin-3-yl]carbamic acid, 1,1-dimethylethyl ester (540 mg, 2.4 mmol) and methyl isocyanate (200 mg, 3.51 mmol) in toluene (20 mL) in a sealed tube was kept at 90 °C for 5 days, then cooled to rt. After the removal of solvents *in vacuo*, the crude product was purified by column chromatography with 1% MeOH in CH₂Cl₂ to afford 680 mg (100%) of the product as a colorless syrup. LC-MS:285 (M⁺+1). ¹H NMR (CDCl₃): δ 1.37 (m, 1H), 1.42 (s, 9H), 1.63 (m, 1H), 1.96 (m, 3H), 2.08 (m, 1H), 2.91 (d, 3H), 3.18 (dd, 1H), 4.63 (m, 1H), 4.84 (m, 1H), 5.67 (d, 1H), 8.96 (s, 1H).

The following intermediates were prepared in a similar manner:

[(3S)-hexahydro-2-oxo-1-[(2-propenylamino)carbonyl]-1H-azepin-3-yl]-carbamic acid-1,1-dimethylethyl ester. ¹H NMR (CDCl₃) δ 1.39 (m, 1H), 1.42 (s, 9H), 1.6 (m, 1H), 1.9 (m, 2H), 2.11 (m, 1H), 3.23 (m, 1H), 3.97 (br. s, 2H), 4.67 (m, 1H), 4.82 (m, 1H), 5.2 (m, 2H), 5.62 (br. s, 1H), 5.89 (m, 1H), 9.23 (br. s 1H).

[(3S)-1-[(2-chloroethyl)amino]carbonyl]hexahydro-2-oxo-1H-azepin-3-yl] carbamic acid-1,1-dimethylethyl ester. ¹H NMR (CDCl₃): δ 1.38 (m, 1H), 1.44 (s, 9H), 1.62 (m, 1H), 1.9 (m, 3H), 2.1 (m, 1H), 3.22 (m, 1H), 3.63 (m, 5H), 4.7 (m, 1H), 4.81 (m, 1H), 5.68 (br. s, 1H), 9.44 (br., s, 1H).

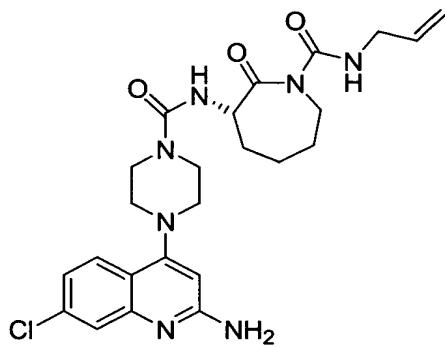
[(3R)-hexahydro-1-[(methylamino)carbonyl]-2-oxo-1H-azepin-3-yl]- carbamic acid-1,1-dimethylethyl ester. ¹H NMR (CDCl₃): δ 1.38 (m, 1H), 1.43(s, 9H), 1.65 (m, 1H), 1.9 (m, 3H), 2.1 (m, 1H), 2.91 (s, 3H), 3.17 (m, 1H), 4.68 (m, 1H), 4.82 (m, 1H), 5.66 (br. s, 1H), 8.97 (br.s, 1H).

Preparation of (3S)-3-[[[4-(2-amino-7-chloro-4-quinoliny)-1-piperazinyl]carbonyl]amino]hexahydro-N-methyl-2-oxo-1H-azepine-1-carboxamide: As described for example 213, [(3R)-Hexahydro-1-[(methylamino)carbonyl]-2-oxo-1H-azepin-3-yl]carbamic acid-1,1-dimethylethyl ester, TFA, triphosgene, NaHCO₃ (sat.),

and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a white powder. LC-MS, 473 (M^+). 1H NMR (400 MHz, DMSO): δ 1.38 (m, 1H), 1.57 (m, 1H), 1.9 (m, 2H), 2.03 (br. d, 1H), 2.19 (s, 1H), 2.85 (d, 3H), 3.08 (m, 4H), 3.21 (m, 1H), 3.63 (m, 4H), 4.83 (m, 1H), 4.85 (m, 1H), 4.95 (s, 2H), 5.85 (d, 1H), 6.11 (s, 1H),
5 7.11 (dd, 1H), 7.56 (d, 1H), 7.67 (d, 1H), 8.91 (q, 1).

Example 260 :

Preparation of (3S)-3-[[[4-(2-Amino-7-chloro-4-quinoliny)-1-piperazinyl]carbonyl]amino]hexahydro-2-oxo-N-(2-propenyl)-1*H*-azepine-1-carboxamide

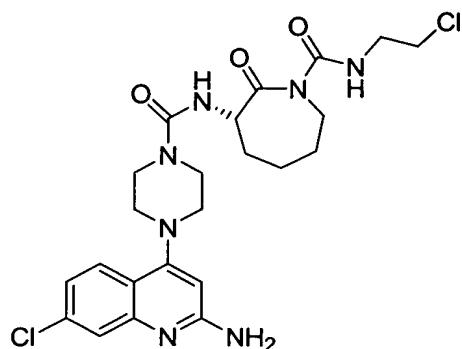


As described for example 78, (3*R*)-3-aminohexahydro-*N*-(2-propenyl)-2-oxo-1*H*-azepine-1-carboxamide obtained by de-protection of [(3*S*)-hexahydro-2-oxo-1-[(2-propenylamino)carbonyl]-1*H*-azepin-3-yl]-carbamic acid, 1,1-dimethylethyl ester with
15 TFA, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to give the product. 1H NMR (DMSO- d_6): δ 1.4 (m, 1H), 1.62 (m, 1H), 1.93 (m, 2H), 2.16 (m, 1H), 3.17 (m, 4H), 3.29 (m, 1H), 3.62 (m, 4H), 3.98 (t, 2H), 4.78 (br., s, 2H), 4.91 (m, 2H), 5.2 (m, 2H), 5.91 (m, 2H), 6.18 (s, 1H), 7.18 (d, 1H), 7.61 (s, 1H), 7.76 (d, 1H), 9.12 (m, 1H).

20

Example 261 :

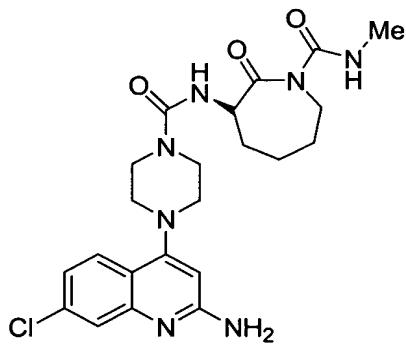
Preparation of (3S)-3-[[[4-(2-Amino-7-chloro-4-quinoliny)-1-piperazinyl]carbonyl]amino]-*N*-(2-chloroethyl)hexahydro-2-oxo-1*H*-azepine-1-carboxamide



As described example 78, (3R)-3-amino-N-(2-chloroethyl)hexahydro-2-oxo-1H-azepine-1-carboxamide obtained by de-protection of [(3S)-1-[[[(2-chloroethyl)amino]carbonyl]hexahydro-2-oxo-1H-azepin-3-yl] carbamic acid-1,1-dimethylethyl ester with TFA, p-nitrophenyl chloroformate, diisopropyl(ethyl)amine, and 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product as a white solid. ^1H NMR (DMSO- d_6): δ 1.41 (m, 1H), 1.63 (m, 1H), 1.79 (m, 1H), 1.96 (m, 3H), 2.17 (m, 1H), 3.16 (m, 4H), 3.34 (dd, 1H), 3.68 (m, 8H), 4.8 (m, 2H), 4.91 (m, 1H), 5.82 (d, 1H), 6.2 (s, 1H), 7.18 (d, 1H), 7.61 (s, 1H), 7.73 (d, 1H), 9.44 (br.s, 1H).

Example 262 :

Preparation of (3R)-3-[[[4-(2-Amino-7-chloro-4-quinoliny)-1-piperazinyl]carbonyl]amino]hexahydro-N-methyl-2-oxo-1H-azepine-1-carboxamide



As described for example 213, [(3R)-hexahydro-1-[(methylamino)carbonyl]-2-oxo-1H-azepin-3-yl]- carbamic acid-1,1-dimethylethyl ester, TFA, K_2CO_3 (sat.) 7-chloro-4-(1-piperazinyl)-2-quinolinamine are reacted to afford the product. ^1H NMR (DMSO- d_6): δ 1.2 (m, 1H), 1.6 (m, 1H), 1.71 (m, 2H), 1.84 (m, 2H), 2.73 (s, 3H), 2.98 (m, 4H), 3.31

(m, 2H), 3.58 (m, 4H), 4.57 (m, 1H), 4.64 (t, 1H), 6.22 (s, 1H), 6.45 (br. s, 1H), 6.78 (d, 1H), 7.11 (d, 1H), 7.37 (s, 1H), 7.76 (d, 1H), 8.84 (m, 1H).

Pyrrolidinylquinolines

5 Example 263

Preparation of 4-[3-(*tert*-Butoxycarbonylamino)pyrrolidin1-yl]-7-chloroquinolin

In a method similar to Example A 4,7-dichloroquinoline (0.49 g, 2.0 mmol), 3-(*tert*-butoxycarbonylamino)pyrrolidine (1.86 g, 10.0 mmol) and 1,4-diazabicyclo[2.2.2]octane (1.12 g, 10.0 mmol) are heated in EtOH for 15 h at reflux. The reaction mixture is concentrated, diluted with EtOAc, washed with water, dried (MgSO_4), and concentrated. The residue is purified by column chromatography with hexane-EtOAc yielding 245 mg of the product as a colorless solid.

^1H NMR (CDCl_3) δ 1.42 (s, 9H), 2.03 (sext, 1H), 2.28 (sext, 1H), 3.50 (dd, 1H), 3.67 (td, 1H), 3.79 (dt, 1H), 3.92 (dd, 1H), 4.39 (br., 1H), 4.81 (br., 1H), 6.41 (d, 1H), 7.29 (dd, 1H), 7.93 (d, 1H), 8.05 (d, 1H), 8.47 (d, 1H).

Example 264

Preparation of 4-[3-(4-Fluorophenylaminocarbonylamino)pyrrolidin1-yl]-7-chloroquinoline

A solution of 4-[3-(*tert*-butoxycarbonylamino)pyrrolidin1-yl]-7-chloroquinoline (214 mg, 0.62 mmol) in CH_2Cl_2 (5 mL) is treated with trifluoroacetic acid (1 mL). After 1 h the reaction mixture is concentrated. The residue is dissolved in CH_2Cl_2 (10 mL), and 4-fluorophenyl isocyanate (70 μL , 0.62 mmol) and triethylamine (1 mL) are added. The reaction mixture is concentrated, and the residue is purified by column chromatography and an aqueous work-up step yielding the title product.

^1H NMR ([D]₆-DMSO) δ 2.07 (sext, 1H), 2.28 (sext, 1H), 3.80 (dd, 1H), 3.97 (m, 2H), 4.20 (dd, 1H), 4.39 (m, 1H), 6.77 (d, 1H), 6.84 (d, 1H), 7.02 (t, 2H), 7.36 (dd, 2H), 7.59 (dd, 1H), 7.89 (d, 1H), 8.45 (d, 1H), 8.48 (d, 1H), 8.62 (s, 1H).

Example 265

Preparation of 4-[3-[*tert*-Butoxycarbonyl(methyl)amino]pyrrolidin1-yl]-7-chloroquinoline

4,7-Dichloroquinoline (0.49 g, 2.0 mmol), 3-[*tert*-butoxycarbonyl(methyl)amino]pyrrolidine (1.96 mL, 10.0 mmol) and 1,4-diazabicyclo[2.2.2]octane (1.12 g, 10.0 mmol) are heated in EtOH for 15 h at reflux. The reaction mixture is concentrated, diluted with EtOAc, washed with water, dried (MgSO_4), and concentrated. The residue is purified by column chromatography with hexane-EtOAc yielding the title product as a colorless solid.

10 ^1H NMR (CDCl_3) δ 1.47 (s, 9H), 2.19 (m, 2H), 2.87 (s, 3H), 3.64 (q, 1H), 3.70 (m, 3H), 4.83 (br., 1H), 6.45 (d, 1H), 7.28 (dd, 1H), 7.94 (d, 1H), 8.07 (d, 1H), 8.48 (d, 1H)

Example 266**Preparation of 4-[3-[4-Fluorophenylaminocarbonyl(methyl)amino]pyrrolidin1-yl]-7-chloroquinoline**

A solution of 4-[3-[*tert*-butoxycarbonyl(methyl)amino]pyrrolidin1-yl]-7-chloroquinoline (285 mg, 0.79 mmol) in CH_2Cl_2 (5 mL) is treated with trifluoroacetic acid (1 mL). After 1 h the reaction mixture is concentrated. The residue is dissolved in CH_2Cl_2 (10 mL), and 20 4-fluorophenyl isocyanate (90 μL , 0.79 mmol) and triethylamine (1 mL) are added. The reaction mixture is concentrated, and the residue is purified by column chromatography and an aqueous work-up step yielding the title product.

1 ^1H NMR ($[\text{D}]_6\text{-DMSO}$) δ 2.18 (m, 1H), 2.25 (m, 1H), 2.97 (s, 3H), 4.00 (m, 4H), 4.97 (m, 1H), 6.80 (d, 1H), 7.06 (t, 2H), 7.44 (dd, 2H), 7.60 (dd, 1H), 7.90 (d, 1H), 8.45 (s, 1H), 8.48 (d, 1H), 8.56 (d, 1H).

Modes of Preparation of Pharmaceutical Compositions**Example 267**

30 This example illustrates the preparation of representative pharmaceutical compositions for oral administration containing a compound of the invention, or a pharmaceutically acceptable salt thereof:

A.	<u>Ingredients</u>	<u>% wt./wt.</u>
	Compound of the invention	20.0%
	Lactose	79.5%
	Magnesium stearate	0.5%

5 The above ingredients are mixed and dispensed into hard-shell gelatin capsules containing 100 mg each, one capsule would approximate a total daily dosage.

B.	<u>Ingredients</u>	<u>% wt./wt.</u>
	Compound of the invention	20.0%
10	Magnesium stearate	0.9%
	Starch	8.6%
	Lactose	69.6%
	PVP (polyvinylpyrrolidone)	0.9%

15 The above ingredients with the exception of the magnesium stearate are combined and granulated using water as a granulating liquid. The formulation is then dried, mixed with the magnesium stearate and formed into tablets with an appropriate tableting machine.

C.	<u>Ingredients</u>	
	Compound of the invention	0.1 g
20	Propylene glycol	20.0 g
	Polyethylene glycol 400	20.0 g
	Polysorbate 80	1.0 g
	Water	q.s. 100 mL

25 The compound of the invention is dissolved in propylene glycol, polyethylene glycol 400 and polysorbate 80. A sufficient quantity of water is then added with stirring to provide 100 mL of the solution, which is filtered and bottled.

D.	<u>Ingredients</u>	<u>% wt./wt.</u>
	Compound of the invention	20.0%
	Peanut Oil	78.0%
30	Span 60	2.0%

The above ingredients are melted, mixed and filled into soft elastic capsules.

E.	<u>Ingredients</u>	<u>% wt./wt.</u>
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Compound of the invention	1.0%
Methyl or carboxymethyl cellulose	2.0%
0.9% saline	q.s. 100 mL

The compound of the invention is dissolved in the cellulose/saline
5 solution, filtered and bottled for use.

Example 268

This example illustrates the preparation of a representative pharmaceutical formulation
for parenteral administration containing a compound of the invention, or a
10 pharmaceutically acceptable salt thereof:

Ingredients

Compound of the invention	0.02 g
Propylene glycol	20.0 g
Polyethylene glycol 400	20.0 g
15 Polysorbate 80	1.0 g
0.9% Saline solution	q.s. 100 mL

The compound of the invention is dissolved in propylene glycol,
polyethylene glycol 400 and polysorbate 80. A sufficient quantity of 0.9% saline
solution is then added with stirring to provide 100 mL of the I.V. solution, which is
20 filtered through a 0.2 m membrane filter and packaged under sterile conditions.

Example 269

This example illustrates the preparation of a representative pharmaceutical
composition in suppository form containing a compound of the invention, or a
25 pharmaceutically acceptable salt thereof:

<u>Ingredients</u>	<u>% wt./wt.</u>
Compound of the invention	1.0%
Polyethylene glycol 1000	74.5%
Polyethylene glycol 4000	24.5%

30 The ingredients are melted together and mixed on a steam bath, and
poured into molds containing 2.5 g total weight.

Example 270

This example illustrates the preparation of a representative pharmaceutical formulation for insufflation containing a compound of the invention, or a pharmaceutically acceptable salt thereof:

	<u>Ingredients</u>	<u>% wt./wt.</u>
5	Micronized compound of the invention	1.0%
	Micronized lactose	99.0%
The ingredients are milled, mixed, and packaged in an insufflator equipped with a dosing pump.		

10 **Example 271**

This example illustrates the preparation of a representative pharmaceutical formulation in nebulized form containing a compound of the invention, or a pharmaceutically acceptable salt thereof:

	<u>Ingredients</u>	<u>% wt./wt.</u>
15	Compound of the invention	0.005%
	Water	89.995%
	Ethanol	10.000%
The compound of the invention is dissolved in ethanol and blended with water. The formulation is then packaged in a nebulizer equipped with a dosing pump.		

20

Example 272

This example illustrates the preparation of a representative pharmaceutical formulation in aerosol form containing a compound of the invention, or a pharmaceutically acceptable salt thereof:

	<u>Ingredients</u>	<u>% wt./wt.</u>
25	Compound of the invention	0.10%
	Propellant 11/12	98.90%
	Oleic acid	1.00%
The compound of the invention is dispersed in oleic acid and the propellants. The resulting mixture is then poured into an aerosol container fitted with a metering valve.		

Example 273

CCR5 Receptor MIP-1a Scintillation Proximity Binding Assay

A) Assay Buffer: 50 mM Hepes, 5 mM MgCl₂, 1 mM CaCl₂, 30 ug/ml bacitracin, 0.1% BSA, pH 7.4.

5

B) Ligand: MIP-1a labeled with I-125 at 20,000 – 25,000 cpm/well. Non specific binding (nsb) was defined as bound cpm in the presence of 100 nM unlabeled MIP-1b.

C) Cells: Human embryonic kidney, (HEK-293) expressing human CCR5 and CD4 pretreated overnight with 5 mM sodium butyrate. Harvest cells with calcium and magnesium free phosphate buffered saline. Cell number is counted with hemacytometer. Cell number per assay point was selected so the total counts per minute (cpm) bound was approximately 10% of the total cpms I-125-MIP-1a added per assay point.

15

D) Beads: Use wheatgerm agglutinin coated scintillation proximity assay beads (sold by Amersham Pharmacia Biotech Inc.) hydrated with the assay buffer for at least an hour before use. Final bead concentration was 0.2 mg beads per well.

E) Scintillation Proximity Assay: 100 ul of assay volume: 60 ul of cell/beads mix (premixed for at least 30 minutes), 20 ul of I-125-MIP-1a, 20 ul of assay buffer for total binding value, or 20 ul of 0.5 uM MIP-1b for nsb, or 20 ul of test compound. Shake the 96 well plates for 30 minutes on an orbital shaker, then let them settle for 30 minutes before reading with a scintillation counter.

25

The preceding examples can be repeated with similar success by substituting the generically or specifically described reactants and/or operating conditions of this invention for those used in the preceding examples.

From the foregoing description, one skilled in the art can easily ascertain the essential characteristics of this invention and, without departing from the spirit and scope thereof, can make various changes and modifications of the invention to adapt it to various usages and conditions.